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

Shriners Hospital of St Louis
Clementia Pharmaceuticals
Courtyard Marriott Earth City

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

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www.mhecoalition.org
Find us on Facebook

MHE Coalition
St Louis Gathering
July 27, 2019
9 am-4 pm CST

Courtyard Marriott Earth City
3101 Rider Trail S, Earth City, MO
(314) 209-1000
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.

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>
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<tr>
<td>9:30-10:30</td>
<td>Clementia Pharmaceuticals</td>
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<tr>
<td>10:30-12:00</td>
<td>Shriners of St Louis</td>
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<tr>
<td>12:00-1:00</td>
<td>Lunch</td>
</tr>
<tr>
<td>1:00-2:00</td>
<td>MHE and ME</td>
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<td></td>
<td>Amy Call, MDiv, MMFT</td>
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<tr>
<td>2:00-3:00</td>
<td>Vocational Rehabilitation</td>
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<td></td>
<td>Matthew Gregg</td>
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<tr>
<td>3:00-4:00</td>
<td>Discussion on Pain, Hospital\Doctor Visit, Hints and Infection control</td>
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</tbody>
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Lunch will be provided

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 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.