ACTION ALERT: Help Support Rare Across America Efforts by Contacting Your Legislators in August!

Rare Across America is now in full swing. **We have nearly 50 PKU patients and advocates throughout the country who are meeting locally with their Members of Congress to advocate for the Medical Nutrition Equity Act during the month of August!** A tremendous thank you to all the participants, we know your efforts will create huge waves of support for the Medical Nutrition Equity Act. For those who are unable to meet with their Members of Congress during August, you can still participate in the action! Call and/or email your Members and urge them to sign on as cosponsor of this bill.

- To send an email to your Representative, click [here](#) and follow the directions provided.
- If you would like to make a phone call, follow this [guide](#).
- Make sure to share with your legislators the Medical Nutrition Equity Act (MNEA) video link: [https://youtu.be/5hW1z1mZOtc](https://youtu.be/5hW1z1mZOtc).

**We would like to make a special ask to all advocates to take action and help buttress the Rare Across America efforts during August. There is GREAT POWER IN NUMBERS and together we truly can make this change for our community. Thank you!**

For questions or to learn more about becoming involved in advocacy, email Kylie Barber at [Kbarber@everylifefoundation.org](mailto:Kbarber@everylifefoundation.org).