

The Honorable Joseph J. Solomon, Jr., Chair House Committee on Corporations The State House 82 Smith Street Providence, RI 02903

## IN SUPPORT OF HOUSE BILL 8041 RELATING TO INSURANCE — PRESCRIPTION DRUG BENEFITS

Dear Chair Solomon and Members of the House Committee on Corportations:

On behalf of the Epilepsy Foundation and our local affiliate, Epilepsy Foundation New England, we write in support of House Bill 8041 regarding the application of cost-sharing amounts toward deductibles or out-of-pocket maximums. This provision would ensure that all people with high prescription drug costs can continue to access needed medications, even if they receive assistance to afford their drugs.

The Epilepsy Foundation is the leading national voluntary health organization that speaks on behalf of more than three million Americans with epilepsy and seizures. Together, we foster the well-being of children and adults affected by seizures through research programs, educational activities, advocacy, and direct services. Epilepsy is a medical condition that produces seizures affecting a variety of mental and physical functions. Approximately 1 in 26 Americans will develop epilepsy at some point in their lifetime. For the majority of people living with the epilepsies, prescription medications are the most common and cost-effective treatment for controlling and/or reducing seizures.

It is beneficial, and at times critical, to our community that individuals are able to receive copayment assistance. Our local affiliate, Epilepsy Foundation New England, provides financial assistance to help pay for the rising costs of medications. Prescription drug manufacturers, and other groups, also offer copayment assistance to individuals who may be struggling to afford their medications for various reasons. If any individuals are not able to afford their medications, even with insurance coverage, and copayment assistance programs play a critical role in ensuring they can gain and maintain seizure control. People living with epilepsy who have their medications switched, or who experience a delay in accessing their medication, are at higher risk for developing breakthrough seizures and related complications including death. Limits on access can also lead to significantly increased medical costs related to preventable seizures, along with lost wages and productivity — not just for the individuals living with epilepsy but also for their family members and communities.

The Epilepsy Foundation and Epilepsy Foundation New England urge you to support H8041and move this legislation forward. Please do not hesitate to contact Dominique Brown, Senior Manager, State Relations & Policy at <a href="mailto:dbrown@efa.org">dbrown@efa.org</a> with any questions.

Sincerely,

Bernice Martin Lee

B. Martin

Chief Executive Officer

**Epilepsy Foundation** 

Susan Linn

President & CEO

Epilepsy Foundation New England

CC:

Representative William W. O'Brien, First Vice Chair Representative Justine A. Caldwell, Second Vice Chair Representative Stephen M. Casey Representative Anthony J. DeSimone Representative Alex S. Finkelman Representative Brian Patrick Kennedy Representative Michelle E. McGaw



The Honorable Joseph J. Solomon, Jr., Chair House Committee on Corporations The State House 82 Smith Street Providence, RI 02903

## IN SUPPORT OF HOUSE BILL 8041 RELATING TO INSURANCE — PRESCRIPTION DRUG BENEFITS

Dear Chair Solomon and Members of the House Committee on Corportations:

On behalf of the Epilepsy Foundation and our local affiliate, Epilepsy Foundation New England, we write in support of House Bill 8041 regarding the application of cost-sharing amounts toward deductibles or out-of-pocket maximums. This provision would ensure that all people with high prescription drug costs can continue to access needed medications, even if they receive assistance to afford their drugs.

The Epilepsy Foundation is the leading national voluntary health organization that speaks on behalf of more than three million Americans with epilepsy and seizures. Together, we foster the well-being of children and adults affected by seizures through research programs, educational activities, advocacy, and direct services. Epilepsy is a medical condition that produces seizures affecting a variety of mental and physical functions. Approximately 1 in 26 Americans will develop epilepsy at some point in their lifetime. For the majority of people living with the epilepsies, prescription medications are the most common and cost-effective treatment for controlling and/or reducing seizures.

It is beneficial, and at times critical, to our community that individuals are able to receive copayment assistance. Our local affiliate, Epilepsy Foundation New England, provides financial assistance to help pay for the rising costs of medications. Prescription drug manufacturers, and other groups, also offer copayment assistance to individuals who may be struggling to afford their medications for various reasons. If any individuals are not able to afford their medications, even with insurance coverage, and copayment assistance programs play a critical role in ensuring they can gain and maintain seizure control. People living with epilepsy who have their medications switched, or who experience a delay in accessing their medication, are at higher risk for developing breakthrough seizures and related complications including death. Limits on access can also lead to significantly increased medical costs related to preventable seizures, along with lost wages and productivity — not just for the individuals living with epilepsy but also for their family members and communities.

The Epilepsy Foundation and Epilepsy Foundation New England urge you to support H8041and move this legislation forward. Please do not hesitate to contact Dominique Brown, Senior Manager, State Relations & Policy at <a href="mailto:dbrown@efa.org">dbrown@efa.org</a> with any questions.

Sincerely,

Bernice Martin Lee

B. Martin

Chief Executive Officer

**Epilepsy Foundation** 

Susan Linn

President & CEO

Epilepsy Foundation New England

CC:

Representative William W. O'Brien, First Vice Chair Representative Justine A. Caldwell, Second Vice Chair Representative Stephen M. Casey Representative Anthony J. DeSimone Representative Alex S. Finkelman Representative Brian Patrick Kennedy Representative Michelle E. McGaw



The Honorable Joseph J. Solomon, Jr., Chair House Committee on Corporations The State House 82 Smith Street Providence, RI 02903

## IN SUPPORT OF HOUSE BILL 8041 RELATING TO INSURANCE — PRESCRIPTION DRUG BENEFITS

Dear Chair Solomon and Members of the House Committee on Corportations:

On behalf of the Epilepsy Foundation and our local affiliate, Epilepsy Foundation New England, we write in support of House Bill 8041 regarding the application of cost-sharing amounts toward deductibles or out-of-pocket maximums. This provision would ensure that all people with high prescription drug costs can continue to access needed medications, even if they receive assistance to afford their drugs.

The Epilepsy Foundation is the leading national voluntary health organization that speaks on behalf of more than three million Americans with epilepsy and seizures. Together, we foster the well-being of children and adults affected by seizures through research programs, educational activities, advocacy, and direct services. Epilepsy is a medical condition that produces seizures affecting a variety of mental and physical functions. Approximately 1 in 26 Americans will develop epilepsy at some point in their lifetime. For the majority of people living with the epilepsies, prescription medications are the most common and cost-effective treatment for controlling and/or reducing seizures.

It is beneficial, and at times critical, to our community that individuals are able to receive copayment assistance. Our local affiliate, Epilepsy Foundation New England, provides financial assistance to help pay for the rising costs of medications. Prescription drug manufacturers, and other groups, also offer copayment assistance to individuals who may be struggling to afford their medications for various reasons. If any individuals are not able to afford their medications, even with insurance coverage, and copayment assistance programs play a critical role in ensuring they can gain and maintain seizure control. People living with epilepsy who have their medications switched, or who experience a delay in accessing their medication, are at higher risk for developing breakthrough seizures and related complications including death. Limits on access can also lead to significantly increased medical costs related to preventable seizures, along with lost wages and productivity — not just for the individuals living with epilepsy but also for their family members and communities.

The Epilepsy Foundation and Epilepsy Foundation New England urge you to support H8041and move this legislation forward. Please do not hesitate to contact Dominique Brown, Senior Manager, State Relations & Policy at <a href="mailto:dbrown@efa.org">dbrown@efa.org</a> with any questions.

Sincerely,

Bernice Martin Lee

B. Martin

Chief Executive Officer

**Epilepsy Foundation** 

Susan Linn

President & CEO

Epilepsy Foundation New England

CC:

Representative William W. O'Brien, First Vice Chair Representative Justine A. Caldwell, Second Vice Chair Representative Stephen M. Casey Representative Anthony J. DeSimone Representative Alex S. Finkelman Representative Brian Patrick Kennedy Representative Michelle E. McGaw



The Honorable Joseph J. Solomon, Jr., Chair House Committee on Corporations The State House 82 Smith Street Providence, RI 02903

## IN SUPPORT OF HOUSE BILL 8041 RELATING TO INSURANCE — PRESCRIPTION DRUG BENEFITS

Dear Chair Solomon and Members of the House Committee on Corportations:

On behalf of the Epilepsy Foundation and our local affiliate, Epilepsy Foundation New England, we write in support of House Bill 8041 regarding the application of cost-sharing amounts toward deductibles or out-of-pocket maximums. This provision would ensure that all people with high prescription drug costs can continue to access needed medications, even if they receive assistance to afford their drugs.

The Epilepsy Foundation is the leading national voluntary health organization that speaks on behalf of more than three million Americans with epilepsy and seizures. Together, we foster the well-being of children and adults affected by seizures through research programs, educational activities, advocacy, and direct services. Epilepsy is a medical condition that produces seizures affecting a variety of mental and physical functions. Approximately 1 in 26 Americans will develop epilepsy at some point in their lifetime. For the majority of people living with the epilepsies, prescription medications are the most common and cost-effective treatment for controlling and/or reducing seizures.

It is beneficial, and at times critical, to our community that individuals are able to receive copayment assistance. Our local affiliate, Epilepsy Foundation New England, provides financial assistance to help pay for the rising costs of medications. Prescription drug manufacturers, and other groups, also offer copayment assistance to individuals who may be struggling to afford their medications for various reasons. If any individuals are not able to afford their medications, even with insurance coverage, and copayment assistance programs play a critical role in ensuring they can gain and maintain seizure control. People living with epilepsy who have their medications switched, or who experience a delay in accessing their medication, are at higher risk for developing breakthrough seizures and related complications including death. Limits on access can also lead to significantly increased medical costs related to preventable seizures, along with lost wages and productivity — not just for the individuals living with epilepsy but also for their family members and communities.

The Epilepsy Foundation and Epilepsy Foundation New England urge you to support H8041and move this legislation forward. Please do not hesitate to contact Dominique Brown, Senior Manager, State Relations & Policy at <a href="mailto:dbrown@efa.org">dbrown@efa.org</a> with any questions.

Sincerely,

Bernice Martin Lee

B. Martin

Chief Executive Officer

**Epilepsy Foundation** 

Susan Linn

President & CEO

Epilepsy Foundation New England

CC:

Representative William W. O'Brien, First Vice Chair Representative Justine A. Caldwell, Second Vice Chair Representative Stephen M. Casey Representative Anthony J. DeSimone Representative Alex S. Finkelman Representative Brian Patrick Kennedy Representative Michelle E. McGaw