

SCHEDULE "A"

COMPENSATION GRID

Injury Tier	Required Documentation	Compensation
<p>Tier I – Symptoms lasting up to 48 hours. Following the consumption of a Recalled Product, the Primary Claimant:</p> <ol style="list-style-type: none"> 1. developed symptoms consistent with <i>Listeriosis</i> (i.e., symptoms developed at least 48 hours after consumption of a Recalled Product and no more than 70 days after consumption of a Recalled Product, and included vomiting, nausea, fever, cramps, diarrhea, constipation, muscle aches, headache, neck stiffness, confusion, loss of balance), and/or 2. developed symptoms consistent with a psychological disorder caused by exposure to the Recalled Product (i.e., a new diagnosis of a medically recognized psychological disorder following consumption of a Recalled Product, with symptoms that are more acute than upset, disgust, anxiety, insomnia or agitation), and <p>developed symptoms that lasted up to 48 hours.</p>	<ol style="list-style-type: none"> 3. If the Primary Claimant was the purchaser of the consumed Recalled Product, proof of purchase, retained packaging, or a declaration attesting to purchase if these documents are not available; and 4. A declaration attesting to consumption of the Recalled Product and physical illness consistent with <i>Listeriosis</i> or psychological disorder caused by exposure to the Recalled Product. 	<p>\$400 per Primary Claimant (single payment to the Primary Claimant, inclusive of all of the Primary Claimant’s claim).</p>
<p>Tier II – Symptoms lasting more than 48 hours and up to one week. Following the consumption of a Recalled Product, the Primary Claimant:</p> <ol style="list-style-type: none"> 1. developed symptoms consistent with <i>Listeriosis</i> (i.e., symptoms developed at least 48 hours after consumption of a Recalled Product and no more than 70 days after consumption of a Recalled Product, and included nausea, fever, cramps, diarrhea, constipation, muscle aches, 	<ol style="list-style-type: none"> 3. If the Primary Claimant was the purchaser of the consumed Recalled Product, proof of purchase, retained packaging, or declaration attesting to purchase if these documents are not available; and 4. A declaration attesting to consumption of the Recalled Product and physical illness consistent with <i>Listeriosis</i> or psychological disorder caused by exposure to the Recalled Product. 	<p>\$1,500 per Primary Claimant (single payment to the Primary Claimant, inclusive of all of the Primary Claimant’s claim).</p>

<p>headache, neck stiffness, confusion, loss of balance), and/or</p> <p>2. developed symptoms consistent with a psychological disorder caused by exposure to the Recalled Product (i.e., a new diagnosis of a medically recognized psychological disorder following consumption of a Recalled Product, with symptoms that are more acute than upset, disgust, anxiety, insomnia or agitation); and</p> <p>developed symptoms that lasted up to one week.</p>		
<p>Tier III – Symptoms lasting more than one week and the Primary Claimant was not hospitalized – Following the consumption of a Recalled Product, the Primary Claimant:</p> <p>1. developed symptoms consistent with <i>Listeriosis</i> (i.e., symptoms developed at least 48 hours after consumption of a Recalled Product and no more than 70 days after consumption of a Recalled Product, and included fever, cramps, diarrhea, constipation, muscle aches, headache, neck stiffness, confusion, loss of balance), and/or</p> <p>2. developed symptoms consistent with <i>Listeriosis</i> (e.g., symptoms developed at least 48 hours after consumption of a Recalled Product and no more than 70 days after consumption of a Recalled Product, fever, cramps, diarrhea, constipation, muscle aches, headache, neck stiffness, confusion, loss of balance) which caused a miscarriage, or</p>	<p>4. If the Primary Claimant was the purchaser of the consumed Recalled Product, proof of purchase, retained packaging, or declaration attesting to purchase if these documents are not available;</p> <p>5. A declaration attesting to consumption of the Recalled Product; and</p> <p>6. Contemporaneous medical reports or records contains a diagnosis of <i>Listeriosis</i> or evidence of symptoms consistent with <i>Listeriosis</i> or evidence of psychological disorder caused by exposure to the Recalled Product.</p>	<p>\$7,000 per Primary Claimant (single payment to the Primary Claimant, inclusive of all of the Primary Claimant’s claim).</p> <p>Subrogated provincial insurer payments.</p>

<p>3. developed symptoms consistent with a psychological disorder caused by exposure to the Recalled Product (i.e., a new diagnosis of a medically recognized psychological disorder following consumption of a Recalled Product, with symptoms that are more acute than upset, disgust, anxiety, insomnia or agitation); and developed symptoms that lasted more than a week, and was not hospitalized.</p>		
<p>Tier IV – Symptoms lasting more than one week and the Primary Claimant was hospitalized but did not develop complications or permanent symptoms - Following the consumption of a Recalled Product, the Primary Claimant:</p> <ol style="list-style-type: none"> 1. developed symptoms consistent with <i>Listeriosis</i> (i.e., symptoms developed at least 48 hours after consumption of a Recalled Product and no more than 70 days after consumption of a Recalled Product, and included vomiting, nausea, fever, cramps, diarrhea, constipation, muscle aches, headache, neck stiffness, confusion, loss of balance), but <i>Listeriosis</i> did not cause them to develop complications (i.e., a brain infection and/or blood poisoning) and no permanent symptoms, and/or 2. developed symptoms consistent with <i>Listeriosis</i> (e.g., symptoms developed at least 48 hours after consumption of a Recalled Product and no more than 70 days after consumption of a Recalled Product, fever, cramps, diarrhea, constipation, muscle aches, 	<ol style="list-style-type: none"> 4. If the Primary Claimant was the purchaser of the consumed Recalled Product, proof of purchase, retained packaging, or declaration attesting to purchase if these documents are not available; 5. A declaration attesting to consumption of the Recalled Product; and 6. Contemporaneous medical reports or records: containing diagnosis or evidence that illness was caused by <i>Listeriosis</i>, recording the number of days an individual was hospitalized, containing Whole Genome Sequencing for the <i>Listeriosis</i> (if available), or documentation that is consistent with the individual having symptoms of <i>Listeriosis</i>, or evidence of psychological disorder caused by exposure to the Recalled Product. 	<p>Capped at \$30,000 per claimant</p> <p>\$10,000 per Primary Claimant (single payment to the Primary Claimant, inclusive of all of the Primary Claimant’s claim).</p> <p>\$900/per day of hospitalization.</p> <p>Special damages (which include and are limited to, reimbursement of reasonable and documented costs, which include, and are limited to, medical expenses, and out-of-pocket expenses for transportation)</p> <p>Subrogated provincial insurers payment.</p>

<p>headache, neck stiffness, confusion, loss of balance) which caused a miscarriage, or</p> <p>3. developed symptoms consistent with a psychological disorder caused by exposure to the Recalled Product (i.e., a new diagnosis of a medically recognized psychological disorder following consumption of a Recalled Product, with symptoms that are more acute than upset, disgust, anxiety, insomnia or agitation) but without permanent symptoms; and</p> <p>developed symptoms that lasted more than a week; and</p> <p>was hospitalized.</p>		
<p>Tier V - Symptoms lasting more than one week where the Primary Claimant was hospitalized and developed severe complications and/or permanent symptoms - Following the consumption of a Recalled Product, the Primary Claimant:</p> <p>1. developed symptoms consistent with <i>Listeriosis</i> (i.e., symptoms developed at least 48 hours after consumption of a Recalled Product and no more than 70 days after consumption of a Recalled Product, and included vomiting, nausea, fever, cramps, diarrhea, constipation, muscle aches, headache, neck stiffness, confusion, loss of balance), and <i>Listeriosis</i> caused the Primary Claimant to develop severe complications (i.e., a secondary infection affecting the central nervous system [i.e., meningoenephalitis or cerebritis, rhombencephalitis, brain abscess, and/or septic shock], and/or focal infections [i.e.,</p>	<p>4. If the Primary Claimant was the purchaser of the consumed Recalled Product, proof of purchase, retained packaging, or declaration attesting to purchase if these documents are not available;</p> <p>5. A declaration attesting to consumption of the Recalled Product; and</p> <p>6. Contemporaneous medical reports or records: containing diagnosis or evidence that illness and secondary illness were caused by <i>Listeriosis</i>, recording the number of days an individual was hospitalized, containing Whole Genome Sequencing for the <i>Listeriosis</i> (if available), or documentation that is consistent with the individual having symptoms of <i>Listeriosis</i>, or evidence of psychological disorder caused by exposure to the Recalled Product.</p>	<p>Capped at \$150,000 per claimant</p> <p>\$30,000 (single payment to the Primary Claimant, inclusive of all of the Primary Claimant's claim).</p> <p>\$900/per day of hospitalization.</p> <p>Subrogated provincial insurers payment.</p> <p>Special damages (which include and are limited to, reimbursement of reasonable and documented costs, which include, and are limited to, medical expenses, and out-of-pocket expenses for transportation).</p> <p>Up to \$70,000 for Family Claimants for Family Claimants (up to \$15,000 for each Family Claimant in relation to the same Class Member). A Family Claimant means a Class Member who is the spouse, child, grandchild, parent, grandparent, brother or sister of a Primary Claimant; resides at the same address as the Primary Claimant, and are not a Primary Claimant.</p>

<p>oculoglandular syndrome, lymphadenitis, pneumonia, empyema, myocarditis, endocarditis, septic arthritis, osteomyelitis, infection of a prosthetic joint, arteritis, infection of a prosthetic graft, spinal or brain abscess, cholecystitis, acute hepatitis or peritonitis]) or caused permanent symptoms, and/or</p> <p>2. developed symptoms consistent with <i>Listeriosis</i> (e.g., symptoms developed at least 48 hours after consumption of a Recalled Product and no more than 70 days after consumption of a Recalled Product, fever, cramps, diarrhea, constipation, muscle aches, headache, neck stiffness, confusion, loss of balance) which caused a stillbirth, or</p> <p>3. developed symptoms consistent with a psychological disorder caused by exposure to the Recalled Product (i.e., a new or significant change in diagnosis of a psychological disorder following consumption of a Recalled Product) but without permanent symptoms; and</p> <p>developed symptoms that lasted more than a week; and</p> <p>was hospitalized.</p>	<p>7. Proof of residency and cohabitation of the Family Claimant and the Primary Claimant for each Family Claimant over 18 years old, or, proof of residency and cohabitation of the Family Claimant's guardian attesting to the cohabitation of the Family Claimant and the Primary Claimant for each Family Claimant over 18 years old.</p>	
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<p>Tier VI – Symptoms resulting in death - Following the consumption of a Recalled Product, the Primary Claimant developed symptoms consistent with <i>Listeriosis</i> (i.e., symptoms developed at least 48 hours after consumption of a Recalled Product and no more than 70 days after consumption of a Recalled Product, and included vomiting, nausea, fever, cramps, diarrhea, constipation, muscle aches, headache, neck stiffness, confusion, loss of balance), resulting in death.</p>	<ol style="list-style-type: none"> 1. If the Primary Claimant was the purchaser of the consumed Recalled Product, proof of purchase, retained packaging, or declaration from the estate attesting to purchase if these documents are not available; 2. A declaration from the estate attesting to consumption of the Recalled Product; 3. Contemporaneous medical reports or records with medical conclusions or evidence that the death was caused by <i>Listeriosis</i>, containing Whole Genome Sequencing for the <i>Listeriosis</i> (if available), or documentation that demonstrates the individual had symptoms consistent with <i>Listeriosis</i>, or evidence of psychological disorder caused by exposure to the Recalled Product. 4. Proof of residency and cohabitation of the Family Claimant and the Primary Claimant for each Family Claimant over 18 years old, or, proof of residency and cohabitation of the Family Claimant's guardian attesting to the cohabitation of the Family Claimant and the Primary Claimant for each Family Claimant over 18 years old. 5. Invoices and receipts related to all funeral expenses. 	<p>Capped at \$300,000 per claimant</p> <p>\$150,000 (single payment to the Primary Claimant, inclusive of all of the Primary Claimant's claim).</p> <p>\$900/per day of hospitalization.</p> <p>Special damages (which include and are limited to, reimbursement of reasonable and documented costs, which include, and are limited to, medical expenses, out-of-pocket expenses for transportation and up to \$19,612.30 for funeral expenses)</p> <p>\$105,000 for Family Claimants (up to \$15,000 for each Family Claimant in relation to the same Class Member). A Family Claimant means a Class Member who is the spouse, child, grandchild, parent, grandparent, brother or sister of a Primary Claimant; resides at the same address as the Primary Claimant, and are not a Primary Claimant.</p> <p>Subrogated provincial insurers payment.</p>
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