Mycotoxin Panel Report Form

Patient: [Name]

Patient Date of Birth: [Date]

Date of Service: [Date]

Date of Report: [Date]

Ordering Physician: Dr. Anna Davis-DirectLabs

Specimen: Urine

Procedure:
- **TYPE:** Aflatoxin (Procedure by ELISA).
- **TYPE:** Ochratoxin (Procedure by ELISA).
- **TYPE:** Trichothecenes (Procedure by ELISA).

Test Results:

<table>
<thead>
<tr>
<th>Code</th>
<th>Test</th>
<th>Specimen</th>
<th>Value</th>
<th>Result</th>
<th>Negative if less than</th>
<th>Equivocal if between</th>
<th>Positive if greater or equal</th>
</tr>
</thead>
<tbody>
<tr>
<td>E8501</td>
<td>Ochratoxin</td>
<td>Urine</td>
<td>0 ppb</td>
<td>Negative</td>
<td>1.8 ppb</td>
<td>1.8-2.0 ppb</td>
<td>2.0 ppb</td>
</tr>
<tr>
<td>E8502</td>
<td>Aflatoxin</td>
<td>Urine</td>
<td>0 ppb</td>
<td>Negative</td>
<td>0.8 ppb</td>
<td>0.8-1.0 ppb</td>
<td>1.0 ppb</td>
</tr>
<tr>
<td>E8503</td>
<td>Trichothecene</td>
<td>Urine</td>
<td>0.05 ppb</td>
<td>Negative</td>
<td>0.18 ppb</td>
<td>0.18-0.2 ppb</td>
<td>0.2 ppb</td>
</tr>
</tbody>
</table>

*Tests such as this should be used only in conjunction with other medically established diagnostic elements (e.g., symptoms, history, clinical impressions, results from other tests, etc). Physicians should use all the information available to them to diagnose and determine appropriate treatment for their patients.*

Director's Initials: **JSS**

Disclaimer: This test was developed and its performance characteristics determined by RealTime Lab. It has not been cleared or approved by the U.S. Food and Drug Administration. The FDA has determined that such clearance or approval is not necessary. This laboratory is certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA-88) as qualified to perform high complexity clinical laboratory testing.