

## New England Biolabs Certificate of Analysis

**Product Name:** *Ph.D.<sup>™</sup>-12 Phage Display Peptide Library Kit*

**Catalog #:** *E8110S*

**Kit Components:** *Ph.D.<sup>™</sup>-12 Phage Display Peptide Library (E8111) — Store at -20°C*  
*-96 gIII Sequencing Primer (20-mer) (S1259) — Store at -20°C*  
*-28 gIII Sequencing Primer (22-mer) (S1258) — Store at -20°C*  
*E. coli K12 ER2738 (E4104) — Store at -80°C*  
*Biotin (N7024) — Store at -20°C*  
*Streptavidin, lyophilized (N7023) — Store at -20°C*

**Lot #:** *0431801*

**Assay Date:** *01/2018*

**Expiration Date:** *01/2020*

**Storage Temp:** *Multi-temperature*

**Specification Version:** *PS-E8110S v1.0*

**Effective Date:** *18 Jun 2018*

Assay Name/Specification (minimum release criteria)	Lot #0431801
<p><b>Absolute Phage Titer</b> - Infection of a mid-log culture of <i>E. coli</i> ER2738 with Ph.D.<sup>™</sup>-12 Phage Display Peptide Library followed by plating, yields <math>\geq 1 \times 10^{13}</math> pfu/ml.</p>	<b>Pass</b>
<p><b>Functional Testing (Panning)</b> - A 100-fold representation of the Ph.D.<sup>™</sup>-12 Phage Display Peptide Library containing approximately <math>10^{11}</math> pfu is diluted in 200 <math>\mu</math>l TBS and panned against 300 ng <math>\beta</math>-endorphin monoclonal antibody. The bound phage is affinity captured using magnetic beads and eluted with 1 ml of 0.2M Glycine-HCl, pH 2.2. After three rounds of selection, <math>\geq 75\%</math> of sequences contain a motif related to the known epitope for the antibody.</p>	<b>Pass</b>
<p><b>Phage Contamination (Environmental)</b> - A 1:100 dilution of an overnight culture of <i>E. coli</i> ER2738 was made in 20 ml LB, to which <math>10^5</math> pfu of Ph.D.<sup>™</sup>-12 Phage Display Peptide Library was added. The flask was incubated at 37°C on a rotating shaker for 5 hours. A 1 ml volume of culture was removed and centrifuged. Five microliters (5 <math>\mu</math>l) of phage-containing supernatant was used for three successive rounds of amplification. The final culture supernatant was plated on three LB/IPTG/Xgal plates and then titered. Fewer than 5% clear or white plaques were observed in a minimum of 100 total plaques counted on each plate.</p>	<b>Pass</b>



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<b>Assay Name/Specification</b> (minimum release criteria)	<b>Lot #0431801</b>
<b>Sequence Verification (DNA)</b> - The Ph.D. <sup>™</sup> -12 Phage Display Peptide Library was sequenced using 5'-CCCATGTACCGTAACACTGAGTTTC-3' as a primer to confirm the correct form of the cloned insert on the displayed peptide, X <sub>12</sub> -GGG.	<b>Pass</b>



Authorized by  
Derek Robinson  
18 Jun 2018



Inspected by  
Beth Paschal  
02 Jan 2018

