

**18S + 28S Ribosomal RNA  
From Calf Liver**

Product No. **R0889**

Lot 051K1468  
Store at -70 °C

**PRODUCT SUMMARY**

Suitable for use as a molecular weight marker for formaldehyde agarose gel electrophoresis.

Concentration: 2.0 mg/ml

**STORAGE BUFFER**

10 mM Tris, pH 8.0  
1 mM EDTA

**10X MOPS ELECTROPHORESIS BUFFER**

400 mM MOPS  
100 mM Sodium Acetate  
10 mM EDTA

**LOADING DYE**

50 % (v/v) glycerol  
1 mM EDTA  
0.4 % (w/v) Bromphenol Blue  
0.4 % (w/v) Xylene Cyanol

**SUITABILITY ASSAY**

E. coli 18S + 28S ribosomal RNA (rRNA) sample solutions were prepared for electrophoresis as follows:

1.0 µg 18S + 28S Ribosomal RNA  
2.5 µl 10X Running Buffer  
3.5 µl 37% Formaldehyde  
10 µl Deionized Formamide  
1 µl Ethidium Bromide (0.2 mg/ml)

**Product Information****SUITABILITY ASSAY (continued)**

The above sample solution was incubated at 65 °C for 10 minutes and immediately cooled on ice. 2 µl of loading dye was added and the entire sample solution (1 µg 18S + 28S Ribosomal RNA) was run along with appropriate RNA markers on a denaturing (formaldehyde) agarose gel. Electrophoresis was performed in a submarine-type apparatus at 100 volts for 2 hours in 1X MOPS electrophoresis buffer with buffer recirculation. Two bands were resolved, and the sizes were consistent with 18S and 28S ribosomal RNA.

**Fragment Sizes:**

18S rRNA approx. 2000 bases  
28S rRNA approx. 5300 bases

**References:**

1. Sambrook, J., et al., **Molecular Cloning, A Laboratory Manual**, Cold Spring Harbor Laboratory (1989), p.202.
2. Fasman, G.D., ed., **Practical Handbook of Biochemistry and Molecular Biology**, CRC Press, (1986), p.464.