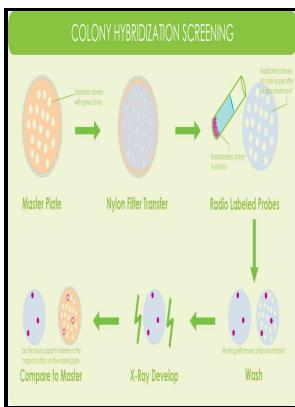


Generation of cDNA libraries - methods and protocols

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Methods in Molecular Biology (Clifton, N.J.) -- 221
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 Notes: Includes bibliographical references and index
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Generation of cDNA Libraries

Positive Control pEXP7-tet control DNA is included to use as a positive control for the BP reaction.

Comparative Analysis of Single

You may screen your cDNA library to identify a specific entry clone and use this entry clone in an LR recombination reaction with a destination vector to generate an expression clone. If your cDNA yield is less than 75 ng, you may pool additional fractions and ethanol precipitate the cDNA. Transformation Control pUC19 plasmid is included to check the transformation efficiency of ElectroMAX DH10B T1 Phage Resistant Cells.

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Design your reverse primer such that it is 5' - T 23N-3' where N is A, C, or G. Do not add scintillation fluid.

5 Steps to Optimal cDNA Synthesis

Use a fresh tip for each fraction to avoid cross-contamination. Contact your safety department for regulations regarding radioactive waste disposal. Thank you for visiting nature.

Generation of cDNA Libraries

Keep reactions at 45°C when adding SuperScript II RT. Place the tube on ice. Generally, spots containing more than 50 ng of cDNA will appear equally stained under UV light.

cDNA Library Construction

Proceed to Collecting Fractions, below. Cite this protocol as: Rederstorff M. CLM includes long-form articles, events listings, publication reviews, new product information and updates, reports of conferences and letters.

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