

pluriBead®

Application example

Minimize artifacts in RNA expression and protein profiling

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Minimize artifacts in RNA expression and protein profiling

Introduction

Current cell separation methods can take up to two hours and need multiple centrifugation steps and temperature shifts.

With the pluriBead® kits, you can isolate specific cell types directly from whole blood. You can work at room temperature (and even at 37°C) without centrifugation and stabilize the RNA and proteins in less than 10 minutes. This will minimize the artifacts in your analysis.

Tested lysis buffers

Trizol®, Qiagen RLT® buffer, Promega lysis buffer, Invitek cell lysis buffer, Ambion lysis buffer, PAXgene® and RNAlater®.

What is different to standard cell separation with pluriBead®?

If you want to use your target cells for RNA expression analysis, you don't need to detach them from the pluriBead® particles. Simply add a suitable lysing buffer and follow the protocol for your downstream experiment.

Use a pluriBead® kit to get started.

	Magnetic separation	pluriBead®
Application in whole blood	+/-	Yes
Specific cell isolation	Yes	Yes
Pre-treatment of whole blood	Yes	No
Time to receive lysed cells	30 - 120 minutes	<10 minutes
Centrifugation steps	1 - 7	0
Isolation of exosomes	No	Yes
Gentle use at room temperature or 37°C	No	Yes

Less than 10 minutes needed for isolated and lysed cells. See video tutorial on our website: www.pluriselect.com

Protocol for pluriBead® cell isolation with subsequent RNA stabilization



Add pluriBead® to your sample



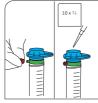
Incubate 4 - 6 minutes



Add sample onto separation device



Wash and discard funnel, wash pluriStrainer®

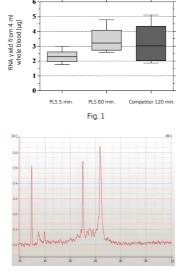


Close Luer-Lock and add lysis buffer



Transfer lysed cells into tube

Figures



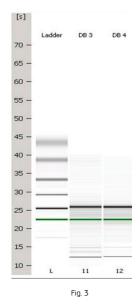


Fig. 1: Comparison of pluriSelect® (PLS) with column technology

Fig. 2: Agilent® RNA integrity analysis

Fig. 2

Fig. 3: Agilent® RNA integrity analysis