



AMPure XP and RNAclean XP Method Overview

The protocol describes the cleanup of either DNA or RNA from enzymatic reactions. Starting input volumes of 5-100 µL from sample is bound by SPRI beads. The bound nucleic acid and SPRI beads are captured with a magnet and the residual solution is transferred to waste. The beads are then washed to remove contaminants and the nucleic acids are eluted from the beads in nuclease-free water.

Extraction Process Workflow

AMPure XP workflow

Aliquot Samples	Add Ampure XP	Wash EtOH	Elution
RNAclean XP workflo	ow.		
Aliquot Samples	Add RNAclean XP	Wash 2x EtOH	Elution

Total Estimated Time

		AMPure XP	RNAclean XP
96 Samples	Hands-on Time	10 min	10 min
	Total Time	19 min	19 min
384 Samples (4 plates)	Hands-on Time	10 min	10 min
	Total Time	43 hr	43 hr

Input Material

Description	Compatible Kit	Storage Temperature
DNA from enzymatic reaction	AMPure XP	4°C— -80°C
RNA from enzymatic reaction	RNAclean XP	4°C80°C

Reagents

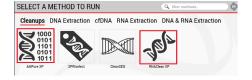
Description	Manufacturer	Storage Temperature
AMPure XP	Beckman Coulter	4°C
RNAclean XP	Beckman Coulter	4°C
100% Ethanol (Molecular Grade)	User	Room Temp
Nuclease-free water (Molecular Grade)	User	Room Temp

Consumables

Description	Supplier	Quantity needed for 1 plate (96 Samples)	Quantity needed for 4 plates (384 Samples)
230μL PIPETTE TIPS, Sterile Filtered	Beckman	1	4
Hard-Shell® Thin-Wall 96-Well Skirted PCR Plates, clear wells	BioRad	2	8
RESERVOIR 96 WELL PYRAMID PP 287ML 25/CS	Agilent	3	3

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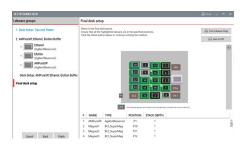
Biomek Method Launcher (BML) organizes methods into useful groups



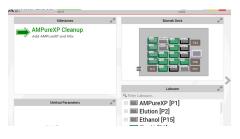
Method Options Selector (MOS) enables flexibility specific to your sample process batch size, process options and workflow customization



Guided Labware Setup (GLS) provides clear instructions to setup the instrument deck with calculated reagent volume and step-by-step instructions to prepare reagents based on options selected



Run Status Screen (RSS) shows progress of run, current activity and time to completion





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