

USB[®] ExoSAP-IT[®] PCR cleanup reagent

The USB ExoSAP-IT method is a unique, patented, one-step enzymatic cleanup of PCR products that is exclusively manufactured and available through Affymetrix.

One-tube, one-step PCR cleanup

Add ExoSAP-IT reagent directly to PCR product

The gold standard in enzymatic PCR cleanup

Over 10,000 publications reference ExoSAP-IT reagent

100% sample recovery

 No loss of PCR products, regardless of the fragment size (Fig. 1)

Eliminates spin columns

Decreases time and expense while increasing yield

Removes contaminating primers & dNTPs

No interference in downstream applications

Simple processing

- Lends itself to robotics
- Replaces beads, filtrations and plates

Scaleable

Treat reaction volumes from 5 µl to 5 L

Stable at 25°C for 8 hours

 Retains full functional activity and is stable at 4°C for one week

Sole source

- Affymetrix is the only manufacturer of USB ExoSAP-IT reagent, ensuring your lab of product consistency and integrity
- Proprietary buffer formulation for superior performance

Green option

 Generates less waste with our single-tube solution to PCR purification vs. columns

ExoSAP-IT reagent

ExoSAP-IT reagent is designed for simple, quick PCR cleanup for downstream applications, such as sequencing or SNP analysis. When PCR amplification is complete, any unconsumed dNTPs and primers remaining in the PCR product mixture will interfere with these methods. ExoSAP-IT cleanup reagent is added directly to the PCR product and incubated at 37°C for 15 minutes. The enzymes are active in the buffer used for PCR, so no buffer exchange is required. After PCR treatment, ExoSAP-IT reagent is inactivated simply by heating to 80°C for 15 minutes.

The ExoSAP-IT method utilizes two hydrolytic enzymes, Exonuclease I and Shrimp Alkaline Phosphatase, together in a specially formulated buffer, to remove unwanted dNTPs and primers from PCR products. Exonuclease I removes residual single-stranded primers and any extraneous single-stranded DNA produced in the PCR. Shrimp Alkaline Phosphatase removes the remaining dNTPs from the PCR mixture which would interfere with the subsequent reactions.

Simple: one step

The method is designed to require a minimum of 'hands-on' time. Enzymatic removal of excess primers and unincorporated nucleotides occurs in one easy step by using ExoSAP-IT reagent in a single tube or microtiter well. Only simple pipette transfers are required, allowing simple automated or manual processing.

100% sample recovery

Use of ExoSAP-IT reagent eliminates all gel or column purifications, sedimentations, filtrations, beads and/or magnetic separations⁽¹⁾. There is 100% recovery of both short and long PCR products with the ExoSAP-IT method.



Single-copy targets were amplified from human genomic DNA. HES-1 (125 bp), numb (455 bp), NRAGE (1.55 kb), and numb (4.6 kb) were loaded on a 1.5% agarose gel before (pre) and after (post) ExoSAP-IT treatment. M is the DNA marker lane. Note that a variety of PCR product sizes can be treated with ExoSAP-IT reagent, even a 125 bp fragment, with no sample loss.

Achieve high data quality

Enzymatic treatment to remove excess primers and nucleotides yields templates which can be easily analyzed. Problems with left-over PCR primers leading to high background bands are virtually eliminated. ExoSAP-IT reagent may be used as an effective cleanup method prior to fluorescent or radioactive DNA sequencing, SNP analysis, or any other application requiring a PCR product free of excess nucleotides and primers.

Storage

Store at -20°C and keep on ice while pipetting. The enzymes are heat-labile but full functional activity is retained after a week of storage at 4°C. Do not store in a frost-free freezer (the temperature rises above 0°C daily).

Composition

Exonuclease I and Shrimp Alkaline Phosphatase in specially formulated buffer.

Rapid PCR product cleanup protocol

The ExoSAP-IT method consists of one pipetting step and two incubations. Just add ExoSAP-IT reagent to the PCR product and within 30 minutes be ready to sequence or do SNP analysis.

ExoSAP-IT PCR Product Cleanup

Product code	Pack size
78250 40 µl	20 reactions
78200 200 µl	100 reactions
78201 1 ml	500 reactions
78202 4 × 1 ml	2,000 reactions
78205 10 ml	5,000 reactions

References:

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- Hanke, M. and Wink, M. (1994) BioTechniques 17, 858-860.
- Mu, J., Duan, J., Makova, K., Joy, D., Huynh, C., Branch, O., Li, W. and Su, X. (2002) Nature 3. 418, 323-326.
- 4 Silva, Jr., W. A., Costa, M. C. R., Valente, V., De Freitas Sousa, J., Pacó-Larson, M. L., Espreafico, E. M., Camargo, S. S., Monteiro, E., De Jesus, A., Holanda, M. A., Zago, M. A., Simpson, A. J. G. and Neto, E. D. (2001) BioTechniques 30, 537-542.
- Werle, E., Scneider C., Renner, M., Völker, M. and Fiehn, W. (1994) Nucleic Acids Res. 22, 4354-4355

Fig. 2. Summary of ExoSAP-IT PCR product treatment.



Related products:

ExoSAP-IT Express PCR Product Cleanup [75001]

PCR cleanup in 5 minutes—ensures quality sequencing results in a fraction of the time.

20 reactions 100 reactions 500 reactions 2,000 reactions 5,000 reactions 480 reactions × 8-tube strip

HT ExoSAP-IT Fast High-Throughput PCR Product Cleanup [78595]

Alternative formulation of the original ExoSAP-IT reagent specifically designed for the unique requirements of high-throughput, automated platforms and multichannel pipettes.

20 reactions

480 reactions × 8-tube strip

5,760 reactions $(12 \times 8$ -tube strips in a tray)

23,040 reactions (48×8 -tube strips in a tray)

1,000 reactions (2 ml)

5,000 reactions (10 ml)

1,920 reactions (96-well plate, 40 µl per well)

11,520 reactions (6×96 -well plate, 40 µl per well)

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