



Liquid Biopsy Research - "We are where you start"

cfKapture™ 21 Kit (200-400 µl)

Manual Revision v1.1

Catalog Nos. CFK-D10-400UL , CFK-D50-400UL

Isolation kit for circulating cell-free DNA from stabilized plasma

PROTOCOL

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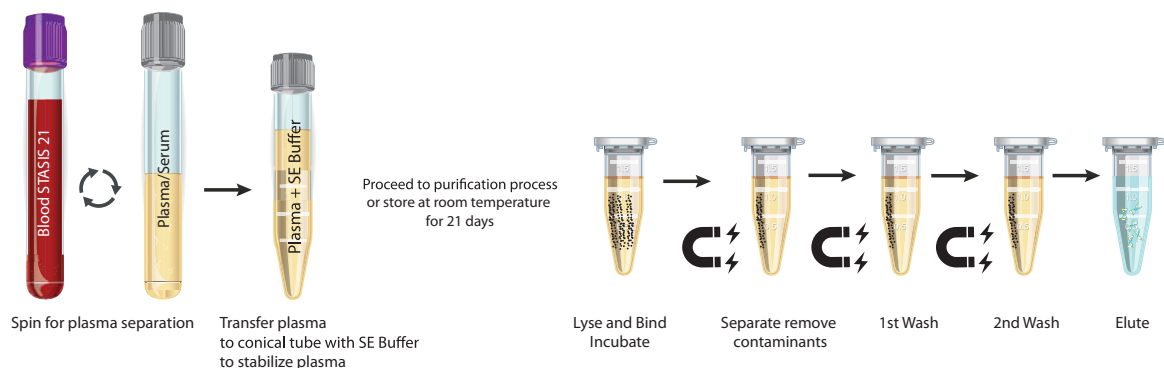
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Product Description

cfKapture™ 21 kit is a system for purification of circulating, cell free nucleic acids from plasma. It is designed for the purification of circulating cell free DNA(ccfDNA) from maternal and cancer patient's whole blood. cfKapture 21 system includes all necessary reagents from the stabilization of separated plasma to the purification of ccfDNA.

The long term room temperature stabilization of plasma during storage or transport is achieved by mixing the plasma sample with our SE buffer allowing for up to 21 day room temperature stabilization of plasma samples. This functionality offers to samples processing laboratories, time flexibility for optimizing their operational efficiency as well as outreach for receiving samples at local and international levels. The isolated cfDNA can be directly used for real time-PCR and DNA library preparation suitable for next generation sequencing. **This kit is designed for research purposes only.**

Process



Kit Contents and Storage

cfKapture 21 Kit Catalog No.	CFK-D10-400UL	CFK-D50-400UL	STORAGE
Number of Preps	10	50	
SE Buffer	21 ml	102 ml	15-25°C
CFL Buffer	6 ml	28 ml	15-25°C
DF Solution	110 µl	520 µl	2-8°C
CFW1 Buffer ¹	8 ml	40 ml	15-25°C
CFW2 Buffer ¹	5 ml	25 ml	15-25°C
Elution Buffer	600 µl	3 ml	15-25°C
Pro K Solution ²	440 µl	2.2 ml	2-8°C
MAG-CFB Particles	220 µl	1.1 ml	2-8°C

¹ Ethanol must be added prior to use. See Preparation of Reagents

Stability

All components are stable for 12 months when stored accordingly.

- Pro K Solution comes in a ready to use solution. Component is stable for 1 year when stored at 15-25°C. For storage longer than 1 year, storage at 2-8°C is recommended.
- During shipment or storage in cool ambient conditions, precipitates may form in some buffers. Dissolve such deposits by warming the solution at 37°C and then gently shaking the buffer.

Safety Information

When working with chemicals, always wear a suitable lab coat, disposable gloves, and protective goggles. For more information, please consult the appropriate material safety data sheets (MSDSs). MSDS can be downloaded from the "Product Resource" tab when viewing the product kit.

Equipments Required

Catalog No.	Description
MBMS-10	Magnetic Separation Device for 2ml microcentrifuge tube

Preparation of Reagents

Prepare the following components for each kit before use:

Catalog No.	Component	Add 100% Ethanol	Storage
CFK-D10-400UL	CFW1 Buffer	10 ml	15-25°C
	CFW2 Buffer	13 ml	15-25°C
Components are stable for 1 year when stored closed at room temperature			

Catalog No.	Component	Add 100% Ethanol	Storage
CFK-D50-400UL	CFW1 Buffer	50 ml	15-25°C
	CFW2 Buffer	65 ml	15-25°C
Components are stable for 1 year when stored closed at room temperature			

CF-Kapture™ 21 Kit : 200 µl plasma sample protocol

Equipment and Reagents to Be Supplied by User

When working with chemicals, always wear a suitable lab coat, disposable gloves, and protective goggles. For more information, please consult the appropriate material safety data sheets (MSDSs) from each product supplier.

- Nuclease-free 1.5 ml microcentrifuge tubes
- 15 ml conical tubes
- magnetic separation device for 1.5 ml microcentrifuge tube (CAT# MBMS-10)
- 100% Ethanol
- Vortex
- Tube rotator
- Water bath, incubator or heat block capable of 60°C

Things to do before starting


- Prepare CFW1 Buffer, CFW2 Buffer according to the “Preparation of Reagents” section on page 2
- and Preheat and warm water bath, incubator or heat block to 60°C
- Warm Elution Buffer to 60°C

Preparing serum/plasma for stabilization

- 1. Add 2ml of SE Buffer to a 15ml conical tube.**
This tube will be used to stabilize the separated plasma from whole blood samples outlined below. 2mL of SE Buffer can stabilize up to 5 mL of plasma.
- 2. Prepare serum/plasma from 9-10ml whole blood:**
 - ⚠ If processing 9-10ml whole blood from a Blood STASIS™ 21 tube (CAT# BS21-CF10-100), centrifugation steps can be performed at room temperature. If not processing from a Blood STASIS™ 21 tube, all centrifugation steps need to be performed at 4°C.**
 - a. Centrifuge whole blood samples at 2,000 g for 15 min.**
 - b. Transfer the plasma to a new 15 ml conical tube.**
 - c. Centrifuge the plasma sample again at 16,000 g for 10 min to remove any residual blood and cell debris.**
 - d. Transfer 4-5 ml of clear plasma to the conical tube containing SE Buffer from step 1 to stabilize the plasma.**
 - e. Stabilized plasma can be stored at room temperature for 21 days or proceed to isolation process.**

Isolation steps

Binding Steps

- 3. Transfer 280 µl stabilized plasma prepared from step 2 to a new 1.5 ml conical tube.**
Note: The 280 µl sample volume factors in the dilution factor.
- 4. Add 40 µl Pro K Solution, and mix well by vortexing at maximum speed for 20 seconds.**
- 5. Incubate sample at 60°C for 10 minutes in a water bath. Mix by inverting the tube once during incubation.**
- 6. Add 280 µl CFL Buffer and mix well by vortexing at maximum speed for 60 seconds, and incubate sample at room temperature for 5 minutes.**
- 7. Add 210 µl 100% ethanol and 20 µl MAG-CFB Particles, and 10 µl of DF Solution. Mix immediately by vortexing at maximum speed for 20 seconds.**
 Complete resuspension of the MAG-CFB Particles is crucial for obtaining purity.
- 8. Incubate sample tube on a tube rotator for 20 minutes at room temperature at 10 rpm.**
Adjust rotator angle to approximately 45 degrees for better mixing.
- 9. Remove the tube from rotator and place on a compatible 1.5 ml magnetic separation device (CAT# MBMS-10) to magnetize the MAG-CFB Particles for 5 minutes or until the magnetic particles are completely cleared from solution.**
- 10. With the tube on the magnetic separation device, remove and discard the cleared supernatant by pipetting.** Do not disturb the attracted beads while aspirating the supernatant.

Wash Steps

- 11. Remove the sample tube off the magnetic separation device.**
- 12. Add 800 µL CFW1 Buffer and resuspend the magnetic particles by vortexing at maximum speed for 1 minute or by pipetting up and down 10 times.**
- 13. Place the 1.5 ml sample tube on a compatible 1.5 ml magnetic separation device (CAT#MBMS-10) to magnetize the MAG-CFB Particles at room temperature for 5 minutes or until the magnetic particles are completely cleared from solution.**
- 14. Remove and discard the cleared supernatant.** Do not disturb the attracted beads while aspirating the supernatant.
- 15. Remove the sample off the magnetic separation device and repeat wash by adding 800µl CFW1 Buffer and resuspend the magnetic particles by vortexing at maximum speed for 1 minute or by pipetting up and down 10 times.**
- 16. Place the 1.5 ml sample tube back on the magnetic separation device to magnetize the MAG-CFB Particles at room temperature for 5 minutes or until the magnetic particles are completely cleared from solution.**
- 17. With the sample still on the magnetic separation device, remove and discard the cleared**

supernatant. Do not disturb the attracted beads while aspirating the supernatant.

18. **Remove the sample tube off the magnetic separation device.**
19. **Add 800 µL CFW2 Buffer and resuspend the magnetic particles by vortexing at maximum speed for 1 minute or by pipetting up and down 10 times.**
20. **Place the 1.5 ml sample tube on the magnetic separation device to magnetize the MAG-CFB Particles at room temperature for 5 minutes or until the magnetic particles are completely cleared from solution.**
21. **With the sample still on the magnetic separation device, remove and discard the cleared supernatant.** Do not disturb the attracted beads while aspirating the supernatant.
22. **Repeat Steps 19-22 for a second CFW2 Buffer wash step.**
23. **With the sample tube still on the magnetic separation device, air dry the MAG-CFB Particles for 10 minutes. Remove any residual liquid with a pipette.**
Note: It is critical to completely remove all liquid from the tube.

Elution Steps

24. **Remove the sample tube containing the MAG-CFB Particles off the magnetic separation device.**
25. **Add 30-50 µl Elution Buffer and completely resuspend the MAG-CFB Particles by vortexing at maximum speed for 10 seconds.**
Note: Heat Elution Buffer at 60°C to improve yield.
26. **Incubate at room temperature for 10 minutes.**
27. **Place the tubes back on the magnetic separation device and wait 5 minutes or until the magnetic particles are completely cleared from elution buffer.**
28. **Transfer the clear supernatant containing the ccfDNA to a new 1.5 ml microcentrifuge tube and store at -20°C.**

cfKapture™ 21 Kit : 400 µl plasma sample

Equipment and Reagents to Be Supplied by User

When working with chemicals, always wear a suitable lab coat, disposable gloves, and protective goggles. For more information, please consult the appropriate material safety data sheets (MSDSs) from each product supplier.

- Nuclease-free 2.0 ml microcentrifuge tubes
- 15 ml conical tubes
- magnetic separation device for 2.0 ml microcentrifuge tube (CAT# MBMS-10)
- 100% Ethanol
- Vortex
- Tube rotator
- Water bath, incubator or heat block capable of 60°C

Things to do before starting

- Prepare CFW1 Buffer, CFW2 Buffer according to the "Preparation of Reagents" section on page 2
- Preheat and warm water bath, incubator or heat block to 60°C
- Warm Elution Buffer to 60°C

Preparing serum/plasma for stabilization

1. Add 2ml of SE Buffer to a 15ml conical tube.

This tube will be used to stabilize the separated plasma from whole blood samples outlined below. 2mL of SE Buffer can stabilize up to 5 mL of plasma.

2. Prepare serum/plasma from whole blood:

⚠ If processing 9-10ml whole blood from a Blood STASIS™ 21 tube (CAT# BS21-CF10-100), centrifugation steps can be performed at room temperature. If not processing from a Blood STASIS™ 21 tube, all centrifugation steps need to be performed at 4°C.

- a. Centrifuge whole blood samples at 2,000 g at room temperature.**
- b. Transfer the plasma to a new 15 ml conical tube.**
- c. Centrifuge the plasma sample again at 16,000 g for 10 min at room temperature to remove any residual blood and cell debris.**
- d. Transfer 4-5 ml of clear plasma to the conical tube containing SE Buffer from step 1 to stabilize the plasma.**
- e. Stabilized plasma can be stored at room temperature for 21 days or proceed to isolation process.**

Isolation steps

Binding Steps

3. **Transfer 560µl (plasma+SE Buffer) from the stabilized plasma sample to a new 2 ml centrifuge tube.**
4. **Add 40 µl Pro K Solution, and mix well by vortexing at maximum speed for 20 seconds.**
5. **Incubate sample at 60°C for 10 minutes in a water bath. Mix by inverting the tube once during incubation.**
6. **Add 560µl CFL Buffer and mix well by vortexing at maximum speed for 60 seconds, and incubate sample at room temperature for 10 minutes.**
7. **Add 420µl 100% ethanol and 20 µl MAG-CFB Particles, and 10 µl of DF Solution. Mix immediately by vortexing at maximum speed for 20 seconds.**

 Complete resuspension of the MAG-CFB Particles is crucial for obtaining purity.

8. **Incubate sample tube on a tube rotator for 20 minutes at room temperature at 10 rpm.** Adjust rotator angle to approximately 45 degrees for better mixing.
9. **Remove the tube from rotator and place on a compatible 2 ml magnetic separation device (CAT# MBMS-10) to magnetize the MAG-CFB Particles for 5 minutes or until the magnetic particles are completely cleared from solution.**
10. **With the tube on the magnetic separation device, remove and discard the cleared supernatant by pipetting.** Do not disturb the attracted beads while aspirating the supernatant.

Wash Steps

11. **Remove the sample tube off the magnetic separation device.**
12. **Add 800 µL CFW1 Buffer and resuspend the magnetic particles by vortexing at maximum speed for 1 minute or by pipetting up and down 10 times.**
13. **Place the 2 ml sample tube on a compatible 2 ml magnetic separation device (CAT#MBMS-10) to magnetize the MAG-CFB Particles at room temperature for 5 minutes or until the magnetic particles are completely cleared from solution.**
14. **Remove and discard the cleared supernatant.** Do not disturb the attracted beads while aspirating the supernatant.
15. **Remove the sample off the magnetic separation device and repeat wash by adding 800µl CFW1 Buffer and resuspend the magnetic particles by vortexing at maximum speed for 1 minute or by pipetting up and down 10 times.**
16. **Place the 2 ml sample tube back on the magnetic separation device to magnetize the MAG-CFB Particles at room temperature for 5 minutes or until the magnetic particles are completely cleared from solution.**
17. **Remove and discard the cleared supernatant.** Do not disturb the attracted beads while aspirating

the supernatant.

- 18. Remove the sample tube off the magnetic separation device.**
- 19. Add 800 µL CFW2 Buffer and resuspend the magnetic particles by vortexing at maximum speed for 1 minute or by pipetting up and down 10 times.**
- 20. Place the 2 ml sample tube on the magnetic separation device to magnetize the MAG-CFB Particles at room temperature for 5 minutes or until the magnetic particles are completely cleared from solution.**
- 21. Remove and discard the cleared supernatant.** Do not disturb the attracted beads while aspirating the supernatant.
- 22. Repeat Steps 19-22 for a second CFW2 Buffer wash step.**
- 23. With the sample tube on the magnetic separation device, air dry the MAG-CFB Particles for 10 minutes. Remove any residual liquid with a pipette.**
Note: It is critical to completely remove all liquid from the tube.

Elution Steps

- 24. Remove the sample tube containing the MAG-CFB Particles off the magnetic separation device.**
- 25. Add 30-50 µl Elution Buffer and completely resuspend the MAG-CFB Particles by vortexing at maximum speed for 10 seconds.**
Note: Heat Elution Buffer at 60°C to improve yield.
- 26. Incubate at room temperature for 10 minutes.**
- 27. Place the tubes back on the magnetic separation device and wait 5 minutes or until the magnetic particles are completely cleared from elution buffer.**
- 28. Transfer the clear supernatant containing the ccfDNA to a new 1.5 ml microcentrifuge tube and store at -20°C.**

Troubleshooting guide

Please use this guide to troubleshoot any problems that may arise. For further assistance, please contact technical support via:

Phone: 1-855-262-4246 (in US), outside US, 1-301-302-0144

Email: support@magbiogenomics.com

Symptoms	Possible Causes	Comments

Ordering Information

Catalog No.	Product	Description	Preps
CFK-D10-400UL	cfKapture 21 Kit (200-400µl) 10 preps	Purification of cell-free DNA (cfDNA) from 200-400 µl STABILIZED plasma	10
CFK-D5-2ML	cfKapture 21 Kit (1-2ml) 5 preps	Purification of cell-free DNA (cfDNA) from 1-2 ml STABILIZED plasma	5
CFK-D5-5ML	cfKapture 21 Kit (3-5ml) 5 preps	Purification of cell-free DNA (cfDNA) from 3-5 ml STABILIZED plasma	5
CFK-D50-400UL	cfKapture 21 Kit (200-400µl) 50 preps	Purification of cell-free DNA (cfDNA) from 200-400 µl STABILIZED plasma	50
CFK-D50-2ML	cfKapture 21 Kit (1-2ml) 50 preps	Purification of cell-free DNA (cfDNA) from 1-2 ml STABILIZED plasma	50
CFK-D50-5ML	cfKapture 21 Kit (3-5 ml) 50 preps	Purification of cell-free DNA (cfDNA) from 3-5 ml STABILIZED plasma	50

Magnetic Separation Devices

Catalog No.	Description
MBMS-31550	15ml and 50ml magnetic stand combo. (3x15ml and 3x50ml)
MBMS-10	MagStip magnetic stand (1.5mL x 10)

Whole blood stabilization tubes

Catalog No.	Product	Description
BS21-CF10-100	Blood STASIS 21-ccfDNA 9 mL (100)	100 tubes: 1 ml Additive, 8 ml blood draw volume
BS21-CF6-100	Blood STASIS 21-ccfDNA 6 mL (100)	100 tubes: 0.6 ml Additive, 5.4 ml blood draw volume
BS21-CF3-200	Blood STASIS 21-ccfDNA 3 mL (200)	200 tubes: 0.35 ml Additive, 2.65 ml blood draw volume



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