# **FITC Amplification Protocol**



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Kit Contents				
Product Code	Description	Volume		
1124-F1-50	Detect F1 (Rabbit anti-FITC)	2 x 20µl		
1124-F2-50	Detect F2 (FITC goat anti rabbit IgG)	2 x 35µl		
1124-DT-25	Detergent (Tween 20)	2 x 1ml		
1124-MD-50	Reagent MD (Antifade + DAPI)	1.25ml		

### **Requirements (not provided)**

Equipment	Reagents
Ethanol cleaned slides	Sodium Chloride
Coverslips	Sodium Citrate
Coplin jars	HCI
Humidified chamber	Formamide
Micro-pipette 1µl, 10µl, 500µl	Clear nail varnish
Pipette 10ml, 20ml	Deionised Distilled water
Vortex	
Parafilm	
Micro-centrifuge	
45°C Water bath	
37°C Incubator	
Fluorescence microscope with a suitable filter set	

#### Approx time:

Preparation 20 min Procedure 70 min

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Solutions to be prepared:	20XSSC 1XSSC 4XSSC Detergent wash solution Stringency wash solution Working Reagent A Working Reagent B			
Solution 20XSSC:	<u>44.1g</u>	NaCl Na Citrate 00ml Deionised Distilled water		
Adjust pH to 7.0 using concentrated HCI (before finalising water volume), aliquot and autoclave. Store at 4°C).				
Solution 1XSSC:		20XSSC Deionised distilled water 1XSSC		
Solution 4XSSC:	<u>400ml</u>	20XSSC Deionised distilled water 4XSSC		
Detergent wash solution:	500ml <u>250µl</u> 500ml	•		
Stringency wash solution:	<u>50ml</u>	Formamide 1XSSC		
100ml Stringency wash solution Stringency wash solution can be reused up to 5 times but should be discarded after 2 months				
Working Reagent A:	6µl	Detection reagent F1		

Working Reagent A:6µlDetection reagent F11244µlDetergent wash solution1250µlWorking Reagent A (F1) (1:200)Incubate in the dark for 5 min. Microcentrifuge at 11.000g for 5 min.

Working Reagent B:	12.5µl	Detection reagent B2	
	<u>1237.5µl</u>	Detergent wash solution	
	1250µl	Working Reagent B (F2) (1:100)	
Incubate in the dark for 5 min. Microcentrifuge at 11.000g for 5 min.			

Note: Ensure all solutions are mixed well.

All solution volumes sufficient for 10 slides

This product is for research use only

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## **Procedure:** Washing

- Pre-warm to 45°C in a water bath at least 30 min before starting: Two Coplin jars of Stringency wash solution (50ml each) Three Coplin jars of Solution 1XSSC (50ml each) One Coplin jar of Detergent wash solution (50ml)
- **Note:** The temperature is important. Check the temperature of the solutions in the Coplin jar and not of the water in the water bath.
  - 2. Take out the slide from the incubator and carefully remove the rubber cement. Place in Solution 1XSSC to remove the coverslip.
- Note: Do not allow to dry.
  - 3. Stringency washes:

Wash slides twice by incubating 5 min each in Stringency wash solution (45°C). Wash slides twice by incubating 5 min each in Solution 1XSSC (45°C). Incubate slide for 4 min in Detergent wash solution. (45°C),

#### **Procedure:** Detection

- 4. Apply 100µl of Working Reagent A onto the slide and cover with Parafilm immediately.
- 5. Incubate slide in a humidified box for 15-20 min at 37°C.
- 6. Remove Parafilm from the slide and wash 3 times for 4 min each time in the Detergent wash solution at room temperature, by emptying and refilling the Coplin jar.
- 7. Apply 100µl of Working Reagent B onto the slide and cover with Parafilm immediately.
- 8. Incubate slide in a humidified box for 15-20 min at 37°C.
- 9. Remove Parafilm from the slide and wash 3 times 4 min in Detergent wash solution at room temperature by emptying and refilling the Coplin jar.
- 10. Drain slide well and mount with 50µl of Reagent MD.
- 11. Apply glass coverslip and seal with nail varnish. Store slides in the dark at 4°C.
- **Note:** You get almost no air bubbles when supplied Reagent MD is applied on the coverslip and the almost dry (but not dried out!) slide is laid face-down on the coverslip.
  - 12. View slides using standard epifluorescence filters for FITC and for counterstain DAPI.