1. Prepare the fallowing mix:

	Final concentrations	Volume
rNTP (Larova)	2mM	80µL
MgCl2	25mM	22,5µL
TrisHCLph8	45mM	22,5µL
DTT	5mM	5µL
Spermidine	1mM	5µL
Pyrophosphatase (Roche)		5µL
T7 purified in the laboratory		5µL
H20		QSP 500µL
Matrice PCR		50µL
25µM blocking strand IDT	(5′-TACTCTGCTATTTTTGCGGGCTTGTA-3′)	Theophylline
	5′-ATTTGGGACTCATCAGCTGG-3′	Guanine

2. Incubate 2h à 37°C

3. Migrate on 8M urea, 12% denaturing polyacrylamide gel using 29:1 acylamide: bis solution (Fisher Scientific) for 1h30 at 200V

4. Reveal by UV shadowing