Transfection with Lipofectamin 2000 (Invitrogen)

Culture	96-	48-	24-	12-	6-	35	60	100	150	T25	T75
Vessel	well	well	well	well	well	mm	mm	mm	mm		
Surface	0.3	0.7	2	4	10	10	20	60	140	25	75
area (cm ²)											
Ratio to	0.2	0.4	1	2	5	5	10	30	70	12.5	37.5
24-well											
Plate											

- ? Cell confluency 80-90% is the best. For each well/dish to be transfected,
- ? dilute LA200 in OptiMEM and DNA in OptiMEM.
- ? Incubate @ RT for 5 min.
- ? Combine the DNA and the LA2000, mix, incubate @ RT for 20 min to allow DNA-lipid complexes to form.
- ? Add the mix directly to each well containing cells and mix gently by rocking the dish.
- ? Return the dish to the CO_2 incubator.
- ? Assay the cells 24-48 h after transfection.

Culture	24-	12- well	6- well,	60 mm	100 mm	T25	T75
Vessel	well		35 mm				
Volume of OpitMEM	50	100	250	500	1500		1800
LA2000, ?1	2	4	10	20	60		75
DNA, ?g	0.5	1	2.5	5	15		20