



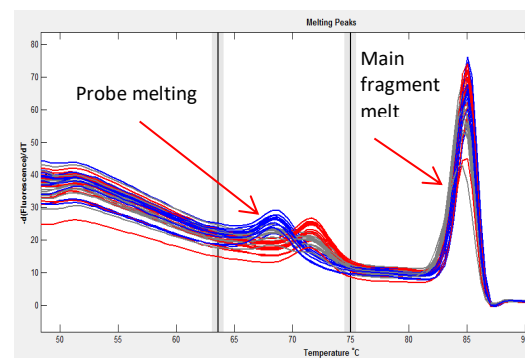
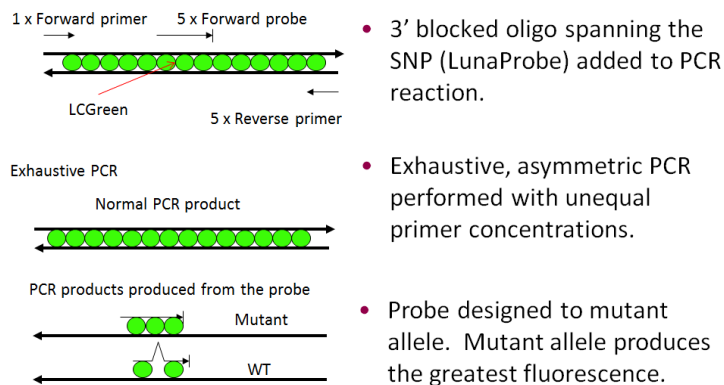
GSP6 Genotyping Strategy

Introduction

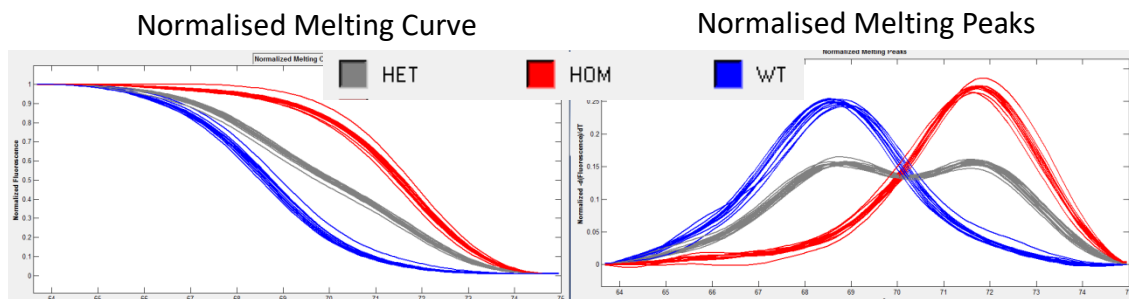
The Idaho Technology LightScanner is a system used to perform high throughput DNA melting analysis. PCR is performed in the presence of the double stranded DNA binding dye LCGreen. After PCR samples are then heated on the LightScanner and the fluorescence emitted by bound LCGreen is monitored. As the DNA melts the LCGreen is released and so the fluorescence decreases until all the DNA has melted and all LCGreen is unbound. There are several different genotyping methods that can be used on the LightScanner.



Unlabelled probe genotyping is used to distinguish between different homozygote samples at a given SNP where scanning analysis may not have enough sensitivity. Here a 3' blocked oligonucleotide (lunaProbe) is designed that sits directly over the SNP. Asymmetric exhaustive PCR is performed using 5 times the amount of probe and opposite primer. This creates two products, one is the full PCR product between the normal primers and the other is the probe that is bound to the opposite strand.



When the products are melted the probe melts at a lower temperature and by focusing analysis on this section hom, het and wt samples can be resolved.





Gasping6

Group: Molecular Embryology
Mutation type: SNP
Mutant allele: A
WT allele: T
Assay Type: LightScanner LunaProbe
Probe direction: Forward

Fragment sequence

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agtacctggtgatcctggggacggtgggtgtgtgggcggatctcaatgtgatgctctctcttcttcatcccac  
ccatattctccgcagCACTTTATGAAAAATGCATGCAGAGAAGAAGCACAAATGCAGTAAGTGCAGTAATTCCTAC  
GGCACCGAGTGGGACCTGAAAAAGACACGAGGAGGATAGTGGCAAGACCTTCCAGTGTACGTGCGGCTGTCCCTAC  
GCCAGCAGAACCCTGCAGTCTCACATCTACCGAACTGGCCACGAGATCCCTGCAGAGCACAGgtaaaacagg  
gagatgocggttgagtagctgtaaatgggacctcctgacaagccagccacaaagggaaacagaaggaacagggggag
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New Primers/Probe sets 5'>3'

Gasping6F2 ATTCCTACGGCACCGAGT
Gasping6R2 GGGACAGCCGCACGTACA
Gasping6PrF2 GACACGAGGAGGATAGTGGCAAGAC

PCR mix

HotShot master mix	5µl
LCGreen	1µl
Gpg6F2 (20ng/µl)	0.1µl
Gpg6R2 (20ng/µl)	0.5µl
Gpg6ProbeF2 (20ng/µl)	0.5µl
DNA (1/10 dil ABI)	2µl
ddH2O	0.9µl

PCR program

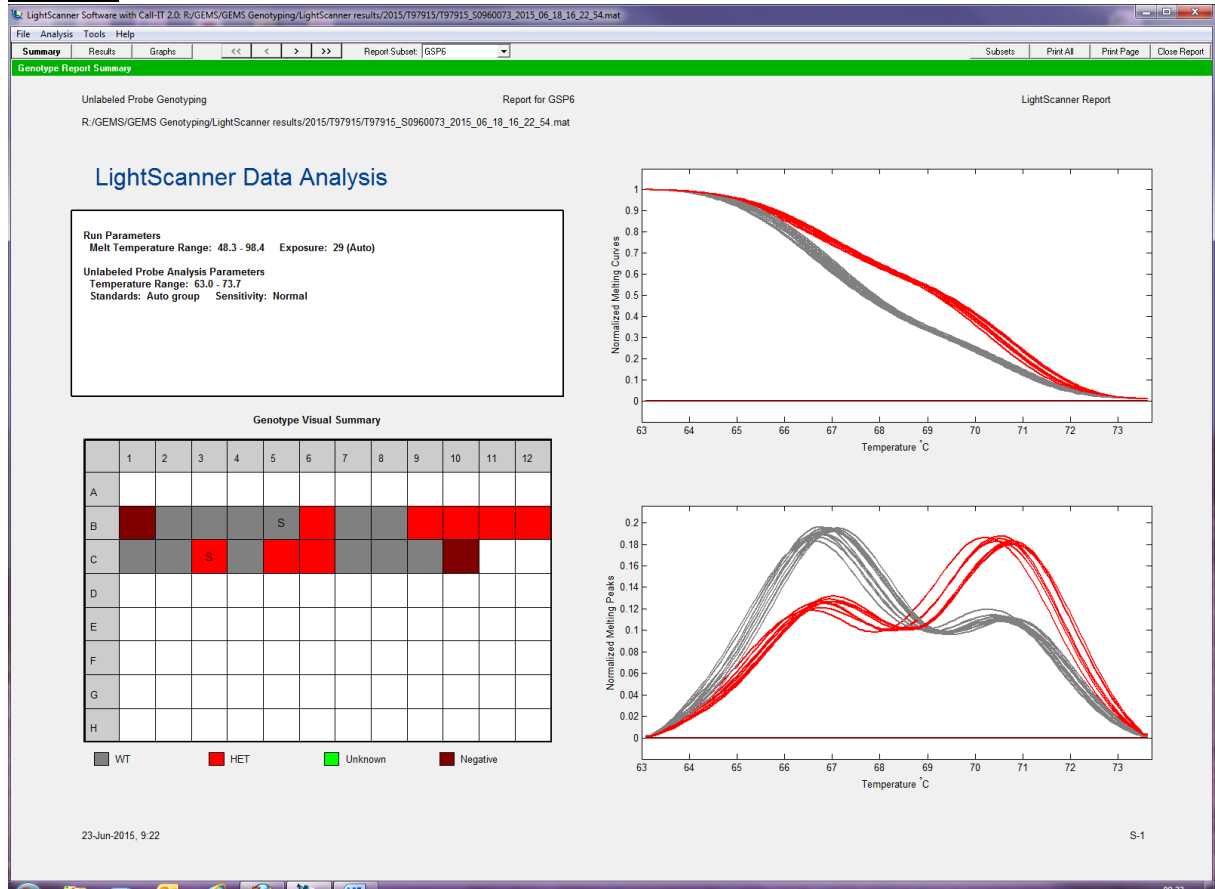
LSGENO60H (annealing temperature 60 °C with hybridisation step)

Control method	Calculated
Lid control mode	Off (no need for heated lid as sample is overlaid with oil)
Lid pressure	Microplate

- 1) 95°C for 2 min
- 2) 95 °C for 30 sec PCR cycle
- 3) 60 °C for 30 sec
- 4) 72 °C for 30 sec
- 5) Cycle, step 2 55 times
- 6) 95 °C for 30 sec Hybridisation
- 7) 25 °C for 30 sec
- 8) 15 °C for 30 sec



Example



HOM

WT HET