# Team3ispA/ispA: PCR

**protocol:** PrimeStar **program:** PG2K55 **thermocycler**:

|  |  |  |  |
| --- | --- | --- | --- |
| **source:**  *label* | *construct* | *concentration* | *location* |
| sF1 dil | sFlexneri | 10uM | Box\_Team3ispA/A3 |
| sF2 dil | sFlexneri | 10uM | Box\_Team3ispA/A4 |

|  |  |  |  |
| --- | --- | --- | --- |
| Sf\*  yP1 dil  yP2 dil  Yp\*  sM1 dil  sM2 dil  Sm  cB1 dil  cB2 dil  Cb\*  pA1 dil  pA2 dil  Pa\* | sFlexneri  yPseudo  yPseudo  yPseudo  sMarc  sMarc  sMarc  citrob  citrob  citrob  pAerug  pAerug  pAerug | miniprep  10uM  10uM  miniprep  10uM  10uM  miniprep  10uM  10uM  miniprep  10uM  10uM  miniprep | genomic\_dnas1/C3  Box\_Team3ispA/B3  Box\_Team3ispA/B4  genomic\_dnas1/F2  Box\_Team3ispA/C3  Box\_Team3ispA/C4  genomic\_dnas1/H5  Box\_Team3ispA/D3  Box\_Team3ispA/D4  genomic\_dnas1/D1  Box\_Team3ispA/E3  Box\_Team3ispA/E4  genomic\_dnas1/E4 |

Tubes marked with \* are actually unlabeled but they should be in the location given on the LabSheet

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **samples:** |  |  |  | |
| *label* | *primer1* | *primer2* | *template product* | |
| ispA1  ispA2  ispA3  ispA4  ispA5 | sF1 dil  yP1 dil  sM1 dil  cB1 dil  pA1 dil | sF2 dil  yP2 dil  sM2 dil  cB2 dil  pA2 dil | sFlexneris sFlexneri\_ispA  yPseudo yPseudo\_ispA  sMarc sMarc\_ispA  citrob citrob\_ispA  pAerug pAerug\_ispA | |
| **Reaction:** |  |  | |  |
|  | 32 uL | ddH2O | | W |
|  | 10 uL | 5X PrimeSTAR GXL Buffer | | B |
|  | 4 uL | PrimeSTAR dNTP Mixture (2.5 mM each) | | D |
|  | 1 uL | 10uM primer 1 | | F |
|  | 1 uL | 10uM primer 2 | | R |
|  | 1 uL | dil20x plasmid template | | T |
|  | 1 uL | PrimeSTAR GXL DNA Polymerase | | P |
| **Notes:** |  |  | |  |

* Do only one thermocycler run for your section
* Never let enzymes warm up! Only take the enzyme cooler out of the freezer when you are actively using it, and only take the tubes out of it when actively dispensing. Hold the enzyme tube by the top of the tube while dispensing and do not place it in a rack.

# Team3ispA/back: PCR

**protocol:** PrimeStar **program:** PGxK55 **thermocycler**:

## source:

*label construct concentration*

*location*

o1 dil o1 10uM Box\_Team3ispA/F3

o2 dil o2 10uM Box\_Team3ispA/F4

T1 pLYC73S dil20x Terp1 /C2

## samples:

*label primer1 primer2 template product*

back1o1 dil o2 dil T1 pLYC73S backbone

|  |  |  |  |
| --- | --- | --- | --- |
| **Reaction:** |  | | |
|  | 32 uL | ddH2O | W |
|  | 10 uL | 5X PrimeSTAR GXL Buffer | B |
|  | 4 uL | PrimeSTAR dNTP Mixture (2.5 mM each) | D |
|  | 1 uL | 10uM primer 1 | F |
|  | 1 uL | 10uM primer 2 | R |
|  | 1 uL | dil20x plasmid template | T |
| **Notes:** | 1 uL | PrimeSTAR GXL DNA Polymerase | P |

* Do only one thermocycler run for your section
* Never let enzymes warm up! Only take the enzyme cooler out of the freezer when you are actively using it, and only take the tubes out of it when actively dispensing. Hold the enzyme tube by the top of the tube while dispensing and do not place it in a rack.