

19. Transfer supernatant to a new tube. Be careful to avoid transferring particles of **Binding Matrix**.
20. Run 2.0 μ l DNA and 2.0 μ l of BBS loading buffer in a 0.8% Agarose gel to check quality / yield.
21. After view the photo of the gel, make aliquots of and identify the by writing the number codes in the top of the tube and in a tape on the tube. Put aliquots in a box and store in the -20.0 C freezer.