

ALTA DIAGNOSTICS, INC. 3123 Research Way Ste 214, Carson City NV 89706 (800) 359-9691 (775) 283-5780 FAX: (775) 283-5787

STABILITY

LIQUID URINE CONTROL **FOR**

MICROSCOPIC & HIGH SPECIFIC GRAVITY

HIGH	35 CELL/HP ± 15 5	SPECIFIC GRAVITY 1.037 1.032 ± .005 1.027	MICROSCOPIC 0 CELL/HP	SPECIFIC GRAVITY 1.037 1.032 ± .005 1.027	PROCEDURE 1. Shake well before using to assure complete mixing of the contents. 2. Remove bottle cap and pour 12 ml into a clean, dry conical centrifuge tube.* 3. Centrifuge for 5 minutes at 2000 rpm. (A lower rpm may be used if this is called for in your laboratory procedure. However, a somewhat lower mean may result!) 4. Remove control from the centrifuge and at this time, if desired, take and record the specific gravity reading by placing a small urinometer in the centrifuge tube or, alternatively, transfer a few drops of the supernate to a refractometer. 5. Pour off and discard all but 0.5 ml of the supernate.		
MEAN 20 C LOW DAY 1 DAY 2 DAY 3 DAY 4 DAY 5 DAY 6 DAY 7 DAY 8 DAY 9 DAY 10 DAY 11 DAY 12 DAY 13 DAY 14 DAY 15 DAY 16 DAY 17 DAY 18 DAY 19 DAY 20	CELL/HP ± 15	1.032 ± .005	0 CELL/HP	1.032 ± .005	complete mixing of the contents. 2. Remove bottle cap and pour 12 ml into a clean, dry conical centrifuge tube.* 3. Centrifuge for 5 minutes at 2000 rpm. (A lower rpm may be used if this is called for in your laboratory procedure. However, a somewhat lower mean may result!) 4. Remove control from the centrifuge and at this time, if desired, take and record the specific gravity reading by placing a small urinometer in the centrifuge tube or, alternatively, transfer a few drops of the supernate to a refractometer. 5. Pour off and discard all but 0.5 ml of the supernate.		
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DAY 8 DAY 9 DAY 10 DAY 11 DAY 12 DAY 13 DAY 14 DAY 15 DAY 16 DAY 17 DAY 18 DAY 19 DAY 20					alternatively, transfer a few drops of the supernate to a refractometer. 5. Pour off and discard all but 0.5 ml of the supernate.		
DAY 9 DAY 10 DAY 11 DAY 12 DAY 13 DAY 14 DAY 15 DAY 16 DAY 17 DAY 18 DAY 19 DAY 20					supernate to a refractometer. 5. Pour off and discard all but 0.5 ml of the supernate.		
DAY 10 DAY 11 DAY 12 DAY 13 DAY 14 DAY 15 DAY 16 DAY 17 DAY 18 DAY 19 DAY 20					supernate.		
DAY 11 DAY 12 DAY 13 DAY 14 DAY 15 DAY 16 DAY 17 DAY 18 DAY 19 DAY 20					supernate.		
DAY 12 DAY 13 DAY 14 DAY 15 DAY 16 DAY 17 DAY 18 DAY 19 DAY 20					6. Resuspend the sediment in the remaining		
DAY 13 DAY 14 DAY 15 DAY 16 DAY 17 DAY 18 DAY 19 DAY 20					6. Resuspend the sediment in the remaining		
DAY 14 DAY 15 DAY 16 DAY 17 DAY 18 DAY 19 DAY 20					0.5 ml of supernate by touching the bottom of the tube to a vortex machine or by flicking the		
DAY 15 DAY 16 DAY 17 DAY 18 DAY 19 DAY 20					bottom of the tube with your finger.		
DAY 16 DAY 17 DAY 18 DAY 19 DAY 20					7. Transfer a drop of the resuspended		
DAY 17 DAY 18 DAY 19 DAY 20					sediment to a clean dry microscope slide and		
DAY 18 DAY 19 DAY 20					cover with a cover slip.		
DAY 19 DAY 20					8. Count and record the <i>average</i> number of cells found in 10 high power fields.		
DAY 20					- · ·		
					9. At the end of the month, add the column of entries for MICROSCOPIC and/or SPECIFIC		
DATZI					GRAVITY and enter the TOTAL at the bottom of the column. Determine the MEAN by		
DAY 22					dividing the TOTAL by the number of days		
DAY 22					the test was run.		
DAY 23					10. Store at 2º - 8ºC. May be stored at room		
DAY 24 DAY 25					temperature once bottle is in use.		
DAY 26					*NOTE:The value range for Alta's Microscopic Control is based on the parameters set forth in the		
DAY 27					above procedure. Laboratories using a procedure		
DAY 28					with different parameters (i.e. volume, rpm and time of centrifugation and amount of supernate		
DAY 29					discarded) should develop their own range of values and mean for the control using their procedure.		
DAY 30					1		
DAY 31					1		
TOTAL					1		
MEAN					1		