

September 04, 2008

Dear Valued Client:

As you may be aware, in recent years there has been a tremendous challenge in our industry to retain trained cytogenetic staff to meet with the testing demand in this growing specialty. Your facility may have experienced delays in turnaround time, in part due to fluctuating staff levels at *Specialty* or other laboratories. While *Specialty* has been fortunate in recent months to have full staffing, we have determined that a new approach to our cytogenetic test offering will ensure a more stable and reliable delivery of test performance. Effective November 4, 2008, *Specialty* will combine its resources with those of our two closest affiliated laboratories. This aggregation will mean that staff from three laboratories will coalesce to provide an enhanced service level. Additional staffing will join our Valencia facility to provide the diagnostic portion of the testing process and the pre-analytical procedures will be performed in the San Juan Capistrano facility. We believe this merger of resources will provide you with quality testing, uninterrupted service levels and better turn around time.

In order to facilitate these changes, we will need to create new test codes for our offerings. Please see the listing below for current offerings and the replacement codes that will be in effect. If you currently order these tests, please make the appropriate changes to your Laboratory Information Systems.

Thank you for choosing *Specialty* and we look forward to your continued support.

For additional information, please visit our Web site at www.specialtylabs.com or contact Client Relations at 800-421-4449.



Christopher Lockhart, M.D.
Laboratory Director

Affected Tests:

Effective November 4, 2008:

- 5880 Acute Lymphoblastic Leukemia by FISH**
Replacement: S51716 - FISH, ALL, Pre-B Panel
- 5874 Acute Myeloid Leukemia/High Grade Myelodysplasia by FISH**
Replacement: S51714 - FISH, MDS/Myeloid Panel, -5/5q-, -7/7q-, +8,20q
- 5840 Acute Promyelocytic Leukemia t(15;17) [FISH]**
Replacement: S51693 - FISH, AML M3, PML/RARA, Translocation 15,17
- 12992 Amniocyte Culture**
Replacement: S51683 - Cell Culture for Possible Additional Prenatal Studies
- 5834 bcr/abl1 Gene Rearrangement (Philadelphia Chromosome) [FISH]**
Replacement: S51686 - FISH, CML/ALL, bcr/abl Translocation 9,22
- 5868 Burkitt Lymphoma t(8;14) by FISH**
Replacement: S51710 - FISH, Burkitts/NHL/ALL, IGH/MYC, t(8;14)
- 5822 Chromosome Analysis Amniotic Fluid**
Replacement: S51687 - Chromosome Analysis, Amniotic Fluid
- 5818 Chromosome Analysis: Products of Conception/Skin Biopsies**
Replacement: S51688 - Chromosome Analysis, Tissue
- 5860 Chronic Lymphocytic Leukemia (CLL) by FISH**
Replacement: S51705 - FISH, B-Cell Chronic Lymphocytic Leukemia (B-CLL) Panel
- 5862 Chronic Myelogenous Leukemia (CML) by FISH**
Replacement: S51686 - FISH, CML/ALL, bcr/abl Translocation 9,22
- 5852 Cytogenetics - inv(16) [FISH]**
Replacement: S51685 - FISH, AML, CBF/MYH11, Inversion 16
- 5854 Cytogenetics - MLL(11q23) [FISH]**
Replacement: S51715 - FISH, MLL (11q23) Gene Rearrangement
- 5850 Cytogenetics - t(8;21) [FISH]**
Replacement: S51682 - FISH, AML, AML1/ETO Translocation 8,21
- 5814 Cytogenetics - Congenital Disorders**
Replacement: S51689 - Chromosome Analysis, Blood
- 5800 Cytogenetics - Hematologic & Neoplastic Disorders**
Replacement: S51690 - Chromosome Analysis, Hematologic Malignancy
- 5825 Cytogenetics - Solid Tumors**
Replacement: S51691 - Chromosome Analysis, Solid Tumor

- 5884 del(4)(q12) by FISH (FIP1L1-PDGFR)**
Replacement: S51711 – FISH, Locus-specific Probe (please specify chromosome and band level if applicable)
- 12990 Fibroblast Culture**
Replacement: S51683 – Cell Culture for Possible Additional Prenatal Studies
- 5843 FISH – Subtelomere Abnormalities [Totalvision]**
Replacement: S51684 – FISH, Subtelomere Screen
- 5892 FISH ALK 2p23**
Replacement: S51706 – FISH, ALCL, ALK, 2p23 Rearrangements
- 5894 FISH BCL6 3q27**
Replacement: S51711 – FISH, Locus-specific Probe (please specify chromosome and band level if applicable)
- 5890 FISH C-MYC 8q24**
Replacement: S51704 – FISH, ALL/NHL, MYC-BA, 8q24 Rearrangement
- 5898 FISH Chromosome 1 Aneuploidy**
Replacement: S51711 – FISH, Locus-specific Probe (please specify chromosome and band level if applicable)
- 5858 FISH Chromosome 18 Aneuploidy**
Replacement: S51711 – FISH, Locus-specific Probe (please specify chromosome and band level if applicable)
- 5856 FISH Chromosome 7 Aneuploidy**
Replacement: S51711 – FISH, Locus-specific Probe (please specify chromosome and band level if applicable)
- 5896 FISH del(13)(q14.3)D13S25**
Replacement: S51711 – FISH, Locus-specific Probe (please specify chromosome and band level if applicable)
- 5869 FISH-20q12 D20S108**
Replacement: S51681 – FISH, Chromosome 20q Deletion
- 5863 FISH-IGH 14q32**
Replacement: S51709 – FISH, B-Cell Malignancy, IGH, 14q32 Rearrangement
- 5865 FISH-MALT1 18q21**
Replacement: S51717 – FISH, MALT Lymphoma, MALT1, 18q21 Rearrangement
- 5861 FISH-p53 17p13.1**
Replacement: S51711 – FISH, Locus-specific Probe (please specify chromosome and band level if applicable)
- 5867 FISH-RB1 13q14**
Replacement: S51711 – FISH, Locus-specific Probe (please specify chromosome and band level if applicable)
- 5885 FISH-Trisomy 13, 18, 21, X & Y [Aneuvysion] (Prenatal)**
Replacement: S51692 – FISH, Prenatal Screen

- 5887 FISH-Trisomy 21**
Replacement: S51711 – FISH, Locus-specific Probe (please specify chromosome and band level if applicable)
- 5872 Follicular Lymphoma t(14;18) by FISH**
Replacement: S51708 – FISH, Follicular Lymphoma, IGH/BCL2, t(14;18)
- 5830 Genetic Diseases [FISH]**
Replacements:
S51697 – FISH, Angelman
S51698 – FISH, Cri du chat
S51699 – FISH, DiGeorge, Velocardiofacial (VCFS)
S51700 – FISH, Miller-Dieker
S51701 – FISH, Smith-Magenis
S51702 – FISH, Wolf-Hirschhorn
S51720 – FISH, SRY/X Centromere
S51721 – FISH, Prader Willi
S51722 – FISH, Williams
S52723 – FISH, X-Linked Ichthyosis Steroid Sulfatase Deficiency
S51724 – FISH, Kallmann
S57103 – FISH, Chromosome-Specific Probe (please specify chromosome and band)
- 5836 HER-2/*neu* [FISH], Breast Cancer**
Replacement: S51696 – FISH, HER-2/*neu*, Paraffin Block
- 5866 MALT Lymphoma by FISH**
Replacement: S51718 – FISH, MALT Lymphoma, MALT1, *rea18q21* w/reflex to API2/MALT1, t(11;18)
- 5870 Mantle Cell Lymphoma t(11;14) by FISH**
Replacement: S51707 – FISH, Mantle Cell Lymphoma, IGH/CCND1, t(11;14)
- 5845 Multiple Myeloma 5 Probes**
Replacement: S51713 – FISH, Myeloma, 13q, 14q, 17p w/reflex to 5,9,15
- 5864 Multiple Myeloma by FISH**
Replacement: S51712 – FISH, Multiple Myeloma, Chromosomes 5,9,15
- 5878 Myelodysplastic Syndromes (MDS), Low Grade by FISH**
Replacement: S51714 – FISH, MDS/Myeloid Panel, -5/5q-, -7/7q-, +8,20q
- 5882 t(12;21) TEL/AML1 by FISH**
Replacement: S51694 – FISH, ALL, TEL/AML1 Translocation 12,21
- 5886 Trisomy 11 by FISH**
Replacement: S51711 – FISH, Locus-specific Probe (please specify chromosome and band level if applicable)
- 5808 Trisomy 8 [FISH]**
Replacement: S51711 – FISH, Locus-specific Probe (please specify chromosome and band level if applicable)
- 5838 X & Y Probe Opposite Sex BM Transplant [FISH]**
Replacement: S51695 – FISH, X/Y, Post Opposite Sex Bone Marrow Transplant

Test Changes:

Effective November 4, 2008:

1833 ER, PR, Ki-67, p53, HER-2/*neu* w/Reflex FISH, Breast Cancer

Component	Specimen Source	(Add)
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1839 ER, PR, Ki-67, HER-2/*neu* w/Reflex FISH, Breast Cancer

Component	Specimen Source	(Add)
Component	ER Breast Interpretation	(Add)
Component	PR Breast Interpretation	(Add)
Component	HER-2/ <i>neu</i> Interpretation	(Add)
Component	Ki-67 Breast Interpretation	(Add)

5846 HER-2/*neu* [IHC] w/Reflex FISH, Breast Cancer

1842 ER, PR, HER-2/*neu* w/Reflex FISH, Breast Cancer

1839 ER, PR, Ki-67, HER-2/*neu* w/Reflex FISH, Breast Cancer

1833 ER, PR, Ki-67, p53, HER-2/*neu* w/Reflex FISH, Breast Cancer

1818 ER, PR, DNA CCA, HER-2/*neu* w/Reflex FISH, Breast Cancer

HER-2/*neu* [IHC] will continue to reflex to FISH when result is 2+, for an additional fee.
Reflexed panel will be S51696 – FISH, HER-2/*neu*, Paraffin Block