Hematology analyzers			
Part 1 of 11 See captodayonline.com/productguides for an interactive version of guide	Abbott Hematology Rick Gooch rick.gooch@abbott.com 5440 Patrick Henry Drive, Santa Clara, CA 95054 800-933-5535 www.abbottdiagnostics.com	Abbott Hematology Rick Gooch rick.gooch@abbott.com 5440 Patrick Henry Drive, Santa Clara, CA 95054 800-933-5535 www.abbottdiagnostics.com	Abbott Hematology Rick Gooch rick.gooch@abbott.com 5440 Patrick Henry Drive, Santa Clara, CA 95054 800-933-5535 www.abbottdiagnostics.com
Name of instrument First year installed in U.S./Outside U.S./No. of units sold in 2011	CELL-DYN Sapphire* 2005/2005/—	CELL-DYN Ruby* 2006/2006/—	CELL-DYN Emerald* 2009/2008/—
No. units installed in U.S./Outside U.S./List price	>190/>800/\$250,000	>550/>2,000/\$185,000	>1,350/>2,000/\$30,000
Test menu: • Chartable (standard menu: WBC, RBC, Hb, Hct, MCV, MCH, MCHC, PLT, %&# neut, mono, lymph, eos, baso):	standard menu (left) plus: MPV, RDW, retic %&#, IRF, NRBC %&#, CD61, CD3T %&#, CD4T %&#, CD8T %&#, 4/8</td><td>standard menu (left) plus: MPV, RDW, retic #& percent</td><td>WBC, RBC, Hb, Hct, MCV, MCH, MCHC, PLT, lymph percent&#, gran percent&#, mid percent&#, RDW, MPV</td></tr><tr><td>• Laboratory • Flags</td><td>— Á band, IG, blast, variant lymph, nvWBC, rstRBC, IR, PLT clmp, ASYM, FP, CD61 agglutination, clot detected during aspiration, short sample</td><td>— NRBC, FWBC, NWBC, RRBC, band, IG, blast, variant lymph, RBC morph., DFLT, MCHC, LRI, URI, LURI, ATYPDEP, high/low interp. message, WBC</td><td>— dispersional data alerts, suspect measurand flags and count invalidation flags</td></tr><tr><td>FDA-cleared tests not clinically released Tests not available but submitted for 510(k) clearance Tests in development</td><td></td><td></td><td></td></tr><tr><td>Tests in development Tests for research use only Tests unique to analyzer</td><td> CD61 for PLTs, CD3/4, CD3/8 (immuno T-cell)</td><td> atypical depolarization flag</td><td></td></tr><tr><td></td><td></td><td></td><td></td></tr><tr><td>Differential method(s) used</td><td>MAPSS (Multi-Angle Polarized Scatter Separation) and three-color fluorescence</td><td>MAPSS (Multi-Angle Polarized Scatter Separation)</td><td>Impedance counting</td></tr><tr><td>Linearity: • WBC count/RBC count • Hemoglobin/Platelet • MCV (fL) or Hct (%) Precision: • WBC count/RBC count • Hemoglobin/Platelet • MCV or Hct Accuracy of automated differential compared with manual differential (per CLSI H-20A2) Interfering substances: • WBC</td><td>$\begin{array}{l} 0.4-250.0\times10^3\;\mu\text{L}/\;0.22-7.50\times10^6\;\mu\text{L}\\ 1.0-24.8\;g/d\text{L}/11.0-2,000.0\times10^3\;\mu\text{L}\\ 37.0-179\;f\text{L}\;(\text{MCV})\\ \leq 2.7\;percent/\leq 1.5\;percent\\ \leq 1.0\;percent/\leq 4.0\;percent\\ \leq 1.0\;percent/\leq 4.0\;percent\\ \leq 1.0\;percent\;(\text{MCV})\\ neut\%\;r=0.942\;slope\;0.947\;y=0.446;\;lym\%\;r=0.936\\ slope=0.943\;y=2.811;\;mon0\%\;r=0.623\;slope=1.057\\ y=0.851;\;eos\%\;r=0.446\;slope=1.024\;y=0.288;\;bas0\%\\ r=0.232\;slope=0.257\;y=0.350\\ \text{PLT}\;clumps,\;neutrophil\;aggregates,\;HbC\;crystals,\\ \end{array}$</td><td>$\begin{array}{l} 0.02-246\times10^{3}/\mu L/0.00-7.50\times10^{6}/\mu L\\ 0.00-25.0\ g/dL/0.00-3,000\times10^{3}/\mu L\\ 58-139\ fL:\ (MCV)\\ 2.4\ percent/1.8\ percent\\ 1.4\ percent/3.8\ percent\\ 0.8\ percent\ (MCV)\\ neut\ percent\ r=0.983,\ slope=0.97,\ y=-1.98;\ lymph\\ r=0.921,\ slope=0.95,\ y=0.94;\ mon\ r=0.711,\\ slope=1.10,\ y=1.93;\ eos\ r=0.952,\ slope=1.04,\\ y=0.01;\ baso\ r=0.146,\ slope=0.18,\ y=1.22\\ fragile\ WBC,\ neutrophil\ aggregates,\ lytic-resistant\\ \end{array}$</td><td>0.4–96.1 K/μL/0.22–7.61 M/μL 3.3–24.6 g/dL/9–1,375 K/uL 5.3–75.6 percent (Hct)/48.8–115 fL (MCV) 3.5% (95% confidence limit)/2.0% (95% confid. limit) 2.1% (95% confidence limit)/6.1% (95% confid. limit) 1.7% Hct (95% confid. limit)/0.8% MCV (95% confid. limit) —</td></tr><tr><td>• RBC</td><td>lyse-resistant RBCs, cryoglobulin, cryofibrinogen, fragmented WBC, NRBCs autoagglutination, cold agglutinins, elevated WBC,</td><td>RBCs, NRBCs, PLT clumps, cryofibrinogen, cryoglobulins elevated WBC, increased numbers of giant PLT,</td><td>proteins, nucleated red cells, platelet clumping, unlysed red cells, clotting, smudge cells, uremia plus immunosuppressants cryoglobulin, cryofibrinogen, giant platelets, high</td></tr><tr><td></td><td>giant PLTs, hemolysis, sm WBC</td><td>autoagglutination, in vitro hemolysis</td><td>white cell count (>50,000 K/µL), autoagglutination, clotting, hemolysis (in vitro), microcytic red cells</td></tr><tr><td>• MCV or Hct</td><td>autoagglutination, cold agglutinins, elevated WBC, giant PLT, hemolysis, hyperglycemia</td><td>MCV: elevated WBC, hyperglycemia, in vitro hemolysis, increased number of giant PLTs</td><td>cryoglobulin, cryofibrinogen, giant platelets, high white cell count (>50,000 K/µL) hyperglycemia (>600 mg/dL), autoagglutination, clotting, hemolysis (in vitro), microcytic red cells, reduced red cell deformability, swollen red cells</td></tr><tr><td>Platelet</td><td>auto and cold agglutinins, cryoglobulins, cryofibrinogen, giant PLT, micro RBCs, PLT clumps, RBC fragments, WBC fragments, PLT satellitism</td><td>WBC fragments, in vitro hemolysis, microcytic RBCs, cryofibrinogen, cryoglobulins, PLT clumping, increased number of giant PLT</td><td>cryoglobulin, cryofibrinogen, hemolysis (in vivo and in vitro), microcytic red cells, red cell inclusions, white cell fragments, clotting, giant platelets, hence in platelet summing platelet outplices</td></tr><tr><td>• Hemoglobin</td><td>lipids>700 mg/dL, WBCs>250 \times 109/L, bilirubin>33 mg/dL, HbC crystals</td><td>elevated WBC, increased plasma substances (triglycerides, bilirubin, in vivo hemolysis), lytic-resistant RBCs</td><td>carboxyhemoglobin (>10 percent), cryoglobulin, cryofibrinogen, hemolysis (in vivo) heparin, high white cell count (>50,000 K/µL), hyperbilirubinemia,</td></tr><tr><td>Interfering substances: Differential</td><td>see WBC</td><td>fragile WBC, neutrophil aggregates, lytic-resistant RBCs, NRBCs, PLT clumps, cryofibrinogen, cryoglobulins, paraproteins</td><td>lipemia, monoclonal proteins platelet aggregates, NRBCs, giant platelets, cryo- globulins, incomplete lysis of RBCs, small lympho- cytes, fibrin clots, shift in WBC cell distrib. due to EDTA anticoagulant equilibration</td></tr><tr><td>Maximum CBCs per hour/Maximum CBCs and differentials per hour</td><td>105/105</td><td>94/94</td><td>60/60</td></tr><tr><td>Minimum specimen volume open/Closed/Sample dead volume closed</td><td>120 µL/120 µL/0.5 mL, 0.3 mL for 10.25 $imes$ 64 mm</td><td>04/04 150 μL/230 μL/1.2 mL</td><td>9.8 µL/—/—</td></tr><tr><td>Microsample capability</td><td>tubes yes</td><td>no</td><td>no</td></tr><tr><td>Prepares microscope slides automatically or flags problems for slide prep Number of automatic slidemakers available/List price</td><td>no —/\$125,000</td><td>no —/\$125,000</td><td>no —</td></tr><tr><td>Archives patient data/Previous patient results incl. with recent results Maximum archived data accessible when system online No. specimens for which numeric results saved in memory at once No. specimens for which histo/cytogram results saved in memory at once Performs delta checks Tags and holds results for followup, confirmatory testing, or rerun Parameters for flags for holding samples defined by user or vendor Scattergram display: cell-specific color Histogram display: color with thresholds User interface can display choice of specimen/result information</td><td>yes/yes 10,000 results 10,000 results 10,000 results yes yes user or vendor yes yes yes</td><td>yes/yes 10,000 results 10,000 results no yes user or vendor yes yes yes</td><td>yes/no 60,000 on USB and 1,500 results on internal memory 60,000 on USB and 1,500 results on internal memory 60,000 on USB and 1,500 results on internal memory no no no no no yes</td></tr><tr><td>LIS interface formats supported</td><td>ASTM 1394</td><td>LIS1/LIS2 CLSI</td><td>proprietary (instrument or vendor specific)</td></tr><tr><td>Information transferred on LIS interface</td><td>numeric and flag results, instrument to LIS; patient demographics, patient orders, LIS to instrument— broadcast; host query for patient demographics and orders</td><td>numeric and flag results, histograms and scat- terplots, instrument to LIS; patient demographics, patient orders, LIS to instrument—broadcast; host</td><td>numeric and flag results, instrument to LIS</td></tr><tr><td>LOINC codes transmitted with all results/Sent in message to LIS/ Listing of machine codes and corresponding LOINC for each test</td><td>no/no/no</td><td>quory for patient demographics and orders no/no/no</td><td>no/no/no</td></tr><tr><td>Interface available or planned to automated specimen-handling system Bar-code symbologies read on specimen tube</td><td>none Codabar, codes 39 and 128, Interleaved 2 of 5</td><td>— Codabar, codes 39 and 128, Interleaved 2 of 5, ISBT</td><td>— Codabar, codes 39 and 128, Interleaved 2 of 5, Chinese post, code 93, EAN8, EAN13, EAN128, IATA, industrial 2 of 5, Italian pharmaceutical, matrix 2 of 5, MSI/Plessey, UK/Plessey, Telepen, TriOptic, S-Code, UPC A, UPC E</td></tr><tr><td>Accommodates bar-code placement per CLSI standard Auto2A</td><td>yes</td><td>yes</td><td>yes</td></tr><tr><td>Time required for maintenance by lab personnel Onboard diagnostics for troubleshooting/Limited to software problems Manufacturer can perform diagnostics via modem</td><td>daily: 30 seconds; weekly: 10 minutes; monthly: 5 minutes yes/no yes</td><td>daily: 30 seconds; weekly: 5 minutes; monthly: 10 minutes yes/no yes</td><td>daily: 3 minutes; monthly: 5 minutes; bi-annually: 10 minutes no/no no</td></tr><tr><td>Distinguishing features (supplied by company) Note: a dash in lieu of an answer means company did not answer question</td><td>four optical and three fluorescent detectors provide multiple scatterplot analysis; 2-D optical platelets prevent interferences; fluorescent analysis of reticulocytes, NRBCs, and three-color monoclonal analysis on routine hematology analyzer; OpenFlow MAb test selections *please see the CELL-DYN Sapphire operator's manual for product</td><td>touch-sensitive screen, all optical technology; onboard maintenance videos; lyse-resistant RBC mode; rules-based result annotations</td><td>small: sample size, reagent volumes used, and physical size; reliable: system averages one service call per year; easy to use: system has touchscreen software with intuitive icons and minimal layers *please see the CELL-DYN Emerald operator's manual for product</td></tr><tr><td>or question is not applicable</td><td>labeling, including warnings, limitations, and precautions</td><td>including warnings, limitations, and precautions</td><td>labeling, including warnings, limitations, and precautions</td></tr></tbody></table>		

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Hematology analyzers

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Part 2 of 11	Beckman Coulter Bill Bailey bdbailey@beckman.com 250 South Kraemer Blvd.	Beckman Coulter Bill Bailey bdbailey@beckman.com 250 South Kraemer Blvd.	Beckman Coulter Bill Bailey bdbailey@beckman.com 250 South Kraemer Blvd.
See captodayonline.com/productguides for an interactive version of guide	Brea, CA 92821 714-961-4440 www.beckmancoulter.com	Brea, CA 92821 714-961-4440 www.beckmancoulter.com	Brea, CA 92821 714-961-4440 www.beckmancoulter.com
Name of instrument First year installed in U.S./Outside U.S./No. of units sold in 2011 No. units installed in U.S./Outside U.S./List price	UniCel DxH 800 2008/2008/— >1,100/>600/\$229,000	LH 1500 Hematology Automation Series 2002/2003/— >50/30/varies	LH 780 2006/2007/— >600/>1,100/LH 780: \$214,500
Test menu: • Chartable (standard menu: WBC, RBC, Hb, Hct, MCV, MCH, MCHC, PLT, %&# neut, mono, lymph, eos, baso):	standard menu (left) plus: RDW-CV, RDW-SD, MPV, retic#, retic%, IRF, MRV, NRBC# and %, body fluids- total nucleated count, and RBC count for synovial, serous, and CSE fluids	standard menu (left) plus: RDW, MPV, retic %&#, IRF, graded RBC morph., NRBC %&#, TNC & RBC on CSF, synovial, and serous fluids</td><td>standard menu (left) plus: RDW, RDW-SD, MPV, retic %&#, IRF, MRV, graded RBC morph., NRBC %&#, TNC and RBC on CSF, synovial, and serous fluids</td></tr><tr><td>• Laboratory • Flags</td><td>definitive, suspect and system messages, user- definable extended decision rules, ISLH consensus rules, user-definable differential sensitivity</td><td>— user-definable age-, gender-, or location-based reference intervals; action and critical limits; user-definable RBC morphology; user-selectable sensitivity for differential, abnormal population suspect messages</td><td>— user-definable age-, gender-, or location-based reference intervals; action and critical limits, user-definable RBC morphology; user-definable sensitivity for differential abnormal populations, suspect and definitive messages</td></tr><tr><td>FDA-cleared tests not clinically released Tests not available but submitted for 510(k) clearance</td><td>_ _</td><td>Ξ.</td><td>-</td></tr><tr><td>Tests in development Tests for research use only</td><td>— high light scatter reticulocytes (HLR% and HLR#), low hemoglobin density (LHD), microcytic anemia factor (MAF), mean sphered cell volume (MSCV), plateletcrit (PCT), platelet distribution width (PDW), reticulocyte distribution width (RDWR-CV and RDWR-SD), red cell</td><td>— MSCV, HLR %&#, PDW, PCT, WBC research population data (RPD); LH 780: MAF, RSF, RDWR-SD, RDWR-CV</td><td>— RSF, MAF, MSCV, HLR %&#, RDWR-CV, RDWR-SD, PDW, PCT, WBC research population data (RPD)</td></tr><tr><td>Tests unique to analyzer</td><td>size factor (RSF), cell population data research parameters</td><td>IVD: NRBCs, body fluids; RUO: MSCV, WBC RPD</td><td>IVD: NRBCs, body fluids, RDW-SD; RUO: MSCV, RSF, MAF, WBC RPD</td></tr><tr><td>Differential method(s) used</td><td>flow cytometric digital analysis using volume, conductivity, and five angles of light scatter, digital signal processing, advanced algorithm applications, high-definition cellular resolution. DataEucian</td><td>Coulter's 3-D VCS biophysical flow cytometry with IntelliKinetics, AccuGate, and AccuFlex technologies</td><td>Coulter's 3-D VCS biophysical flow cytometry with Intellikinetics, AccuGate, and AccuFlex technologies</td></tr><tr><td>Linearity: • WBC count/RBC count • Hemoglobin/Platelet</td><td>0-400/0-8.5 0-25.5/0-3,000</td><td>0-400/0-8.0 0-25/0-3,000</td><td>0-400/0-8.0 0-25/0-3,000</td></tr><tr><td>MCV (fL) or Hct (%) Precision: WBC count/RBC count Hemoglobin/Platelet</td><td>50–150 (MCV) ≤3.0 percent/≤1.5 percent ≤1.5 percent/ ≤3.5 percent</td><td>50–200 (MCV) <1.7 percent/<0.8 percent <0.8 percent/<3.3 percent</td><td>— ≤1.7 percent/≤0.8 percent ≤0.8 percent/≤3.3 percent</td></tr><tr><td>MCV or Hct Accuracy of automated differential compared with manual differential (per CLSI H-20A2)</td><td>\leq1.0 percent NE = ±2.0; LY, MO = ±3.0; EO,BA = ±1.0 (or 10% percent, whichever is greater)</td><td><0.8 percent (MCV) lymph% = ±3.0%, —; neut% = ±3.0%, —; mono% = ±2.0%, eos% = ±1.0%, baso% = ±1.0%</td><td>\leq0.8 percent (MCV) lymph% = ±3.0%, neut% = ±2.0%, mono% = ±3.0%, eos% = ±1.0%, baso% = ±1.0%</td></tr><tr><td>Interfering substances: • WBC</td><td>precipitated elevated proteins, cryoglobulin, fragmented white cells, agglutinated white cells, lyse-resistant red cells, giant platelets, platelet</td><td>unusual RBC abnormalities that resist lysing, NRBCs, fragmented WBC, unlysed particle >35 fL, giant PLT, PLT clumps</td><td>unusual RBC abnormalities that resist lysing, NRBCs, fragmented WBC, unlysed particle >35 fL, giant PLT, PLT clumps</td></tr><tr><td>• RBC</td><td>clumps, unlysed particles >35 fL in size very high WBC count, high concentration of very large platelets, autoagglutinins</td><td>very high WBC, high concentration large PLT, autoagglutinins</td><td>very high WBC, high concentration large PLT, autoagglutinins</td></tr><tr><td>• MCV or Hct</td><td>very high WBC count, high concentration of very large platelets, autoagglutinins</td><td>very high WBC, high concentration large PLT, autoagglutinins</td><td>very high WBC, high concentration large PLT, autoagglutinins (MCV)</td></tr><tr><td>• Platelet</td><td>platelet clumps, white cell fragments, very small red cells, red cell fragments, giant platelets, electric noise</td><td>very small RBCs and WBC fragments</td><td>very small RBCs and WBC fragments</td></tr><tr><td>Hemoglobin Interfering substances: Differential</td><td>severe lipemia, heparin, certain unusual RBC abnormalities that resist lysing elevated triglycerides, precipitated elevated proteins</td><td>very high WBC, severe lipemia, heparin, rare lyse-resistant RBCs high triglycerides may affect lysing</td><td>very high WBC, severe lipemia, heparin, rare lyse-resistant RBCs high triglycerides may affect lysing</td></tr><tr><td>Maximum CBCs per hour/Maximum CBCs and differentials per hour</td><td>>100/>100</td><td>110 per analyzer on automation system/110 per analyzer on automation system</td><td>110/110; 105/100 with SMS</td></tr><tr><td>Minimum specimen volume open/Closed/Sample dead volume closed Microsample capability</td><td>165 µL/165 µL/300–400 µL yes</td><td>200 μL/300 μL, 550 μL with slidemaker/1.0 mL yes</td><td>200 μL/300 μL (550 μL with slidemaker)/1.0 mL yes</td></tr><tr><td>Prepares microscope slides automatically or flags problems for slide prep Number of automatic slidemakers available/List price</td><td>yes —/DxH SMS \$165,000</td><td>yes >850 (U.S.)/\$110,000</td><td>yes >500/\$110,000</td></tr><tr><td>Archives patient data/Previous patient results incl. with recent results Maximum archived data accessible when system online No. specimens for which numeric results saved in memory at once No. specimens for which histo/cytogram results saved in memory at once Performs delta checks Tags and holds results for followup, confirmatory testing, or rerun Parameters for flags for holding samples defined by user or vendor Scattergram display: cell-specific color Histogram display: color with thresholds User interface can display choice of specimen/result information</td><td>yes/yes 40,000 standalone 40,000 standalone 40,000 yes yes yes yes yes yes yes yes</td><td>yes/yes 20,000 samples per instrument 20,000 samples per instrument 20,000 samples per instrument yes yes user or vendor yes yes yes</td><td>yes/yes 20,000 samples 20,000 samples 20,000 samples yes yes user or vendor yes yes yes</td></tr><tr><td>LIS interface formats supported Information transferred on LIS interface</td><td>CLSI LIS01-A2 numeric and flag results, histograms and scatterplots, instrument to LIS; patient demographics, patient orders, LIS to instrument—broadcast; host query for patient demo-</td><td>— numeric and flag results, histograms and scatterplots, instrument to LIS; patient demographics, patient orders, LIS to instrument—</td><td>proprietary numeric and flag results, histograms and scatterplots, instrument to LIS; patient demographics, patient orders, LIS to instrument—</td></tr><tr><td>LOINC codes transmitted with all results/Sent in message to LIS/ Listing of machine codes and corresponding LOINC for each test Interface available or planned to automated specimen-handling system</td><td>graphics and orders (available with release of workcell) no/no/no Beckman Coulter</td><td>proaccast yes/yes/yes Beckman Coulter</td><td>no/no/no Beckman Coulter</td></tr><tr><td>bar-code symbologies read on specimen tube Accommodates bar-code placement per CLSI standard Auto2A</td><td>coaabar, codes 39 and 128, interleaved 2 of 5, NW7 yes</td><td>Locabar, codes 39 and 128, Interleaved 2 of 5, NW7 yes</td><td>uodabar, codes 39 and 128, Interleaved 2 of 5, NW7 yes</td></tr><tr><td>Time required for maintenance by lab personnel</td><td>no routine maintenance. all maintenance</td><td>daily, weekly, monthly and as needed maintenance</td><td>no routine maintenance; only as needed</td></tr><tr><td>Onboard diagnostics for troubleshooting/Limited to software problems Manufacturer can perform diagnostics via modem</td><td>procedures are on as-needed basis yes/no yes</td><td>procedures, nowever time varies by automation line yes/no yes</td><td>yes/no yes</td></tr><tr><td>Distinguishing features (supplied by company) Note: a dash in lieu of an answer means company did not answer question or question is not applicable</td><td>integrated automation with auto repeat-reflex testing based on extended onboard user-defined decision rules; single aspiration pathway negates mode-to- mode comparisons; flow cytometric digital morphol- ogy with five angles of light scatter; separate channel for WBC, NRBC, reticulocyte analysis; digital signal processing, DataFusion; future scalability</td><td>system automatically loads and unloads cassettes, performs reflex and repeat testing, sorts tubes for off-line tests, stores tubes with availability for retrieval for any test type; multiple configurations available; RUO: WBC research population data</td><td>extensive onboard user-defined decision support; extended linearity for WBC and PLT using AccuCount technology; enumeration of NRBCs with every differential; random access-automation ready; integrated slidemaker-slidestainer options; proservice; electronic IQAP; expanded QC module; RUO: WBC research population data</td></tr></tbody></table>	

Tabulation does not represent an endorsement by the College of American Pathologists.

December 2012

Hematology analyzers

Part 3 of 11	Beckman Coulter Bill Bailey bdbailey@beckman.com 250 South Kraemer Blvd.	Beckman Coulter Bill Bailey bdbailey@beckman.com 250 South Kraemer Blvd.	Beckman Coulter Bill Bailey bdbailey@beckman.com 250 South Kraemer Blvd.
See captodayonline.com/productguides for an interactive version of guide	Brea, CA 92821 714-961-4440 www.beckmancoulter.com	Brea, CA 92821 714-961-4440 www.beckmancoulter.com	Brea, CA 92821 714-961-4440 www.beckmancoulter.com
Name of instrument First year installed in U.S./Outside U.S./No. of units sold in 2011	Coulter LH 750 2001/2001/261	Coulter LH 500 2003/2003/198	Coulter HmX 1999 HmX AL/—/— 269 Econocest
No. units installed in U.S./Outside U.S./List price	>2,300/>1,900/\$195,000	>1,500/>1,200/\$145,000	>1,400/>1,800/\$135,000
Test menu: • Chartable (standard menu: WBC, RBC, Hb, Hct, MCV, MCH, MCHC, PLT, %&# neut, mono, lymph, eos, baso):	standard menu (left) plus: RDW, MPV, retic #&%, IRF, MRV, graded RBC morph., NRBC %&#, TNC & RBC on CSF, synovial, and serous fluids</td><td>standard menu (left) plus: retic #, retic %, MRV, IRF, RDW, MPV</td><td>standard menu (left) plus: RDW, MPV, retic #&%, graded RBC morph., IRF, MRV</td></tr><tr><td> Laboratory Flags </td><td>— user-definable age-, gender-, or location-based reference intervals; action and critical limits; user- definable RBC morphology: gradient messages (=+.</td><td>— user-definable age-, gender-, or location-based reference intervals, action and critical limits; user- definable RBC morphology; gradient messages</td><td>— comprehensive high/low, definitive and suspect messages</td></tr><tr><td>FDA-cleared tests not clinically released</td><td>++, +++); user-selectable sensitivity for differential abnormal population suspect messages —</td><td></td><td>_</td></tr><tr><td>Tests not available but submitted for 510(k) clearance Tests in development</td><td></td><td>_</td><td>_</td></tr><tr><td>Tests for research use only</td><td>MSCV, HLR %&#, PDW, PCT, WBC research population data (RPD)</td><td>PCT, PDW, WBC RPD</td><td>PCT, PDW</td></tr><tr><td>Tests unique to analyzer</td><td>IVD: NRBC, body fluids; RUO: MSCV, WBC RPD</td><td>_</td><td>-</td></tr><tr><td>Differential method(s) used</td><td>Coulter's 3-D VCS biophysical flow cytometry with IntelliKinetics, AccuGate, and AccuFlex technologies</td><td>Coulter's 3-D biophysical flow cytometry with AccuGate 500, Reaction Manager technologies</td><td>Coulter's 3-D VCS technology</td></tr><tr><td>Linearity: • WBC count/RBC count • Hemoglobin/Platelet</td><td>0–400/0–8.0 0–25/0–3.000</td><td>0–200/0–7.0 0–25/0–2.000</td><td>0–99.9/0–7.0 0–25/0–999</td></tr><tr><td>MCV (fL) or Hct (%) Precision: • WBC count</td><td></td><td>50-150 (MCV)</td><td>50-150 (MCV)</td></tr><tr><td>Hemoglobin/Platelet</td><td><pre>≤1.7 percent/≤0.8 percent ≤0.8 percent/≤3.3 percent</pre></td><td><pre>≤2.5 percent/≤2.0 percent ≤1.5 percent/≤5.0 percent</pre></td><td><1.5 percent/<5.0 percent</td></tr><tr><td>MCV or Hct Accuracy of automated differential compared with manual</td><td>≤0.8 percent (MCV) lvmph% = ±3.0%, neut% = ±2.0%, mono% =</td><td>≤2 percent (MCV) lvmph= ±1.5 % mean diff., mono= ±1.5 % mean</td><td><2.0 percent (MCV) lymph%= ±3.0%, —; mono%= ±2.0%, —; neut%=</td></tr><tr><td>differential (per CLSI H-20A2)</td><td>$\pm 3.0\%$, eos% = $\pm 1.0\%$, baso% = $\pm 1.0\%$</td><td>diff., neut= $\pm 2.0\%$ mean diff., eos= $\pm 0.5\%$ mean diff. hose $\pm 0.5\%$ mean</td><td>$\pm 3.0\%, -;$</td></tr><tr><td>Interfering substances: • WBC</td><td>unusual RBC abnormalities that resist lysing, NRBC,</td><td>lyse-resistant, nucleated RBCs, fragmented WBCs,</td><td>$105\% = \pm 1.0\%$, —; $1050\% = \pm 1.0\%$, — unusual RBC abnormalities that resist lysing, NRBC,</td></tr><tr><td></td><td>fragmented WBC, unlysed particle >35 fL, giant PLT, PLT clumps</td><td>agglutination WBCs, unlysed particles >35 fL, very large or agg. PLTs, fibrin, cell fragments, or other debris</td><td>fragmented WBC, unlysed particle >35 fL, large PLT</td></tr><tr><td>• RBC</td><td>very high WBC, high concentration large PLT, autoagglutinins</td><td>very high WBC count, many very large PLTs, agglutinin RBCs, RBCs <36 fL, fibrin, cell fragments, or other debris</td><td>very high WBC, high concentration of very large PLT, autoagglutinins</td></tr><tr><td>• MCV or Hct</td><td>MCV and Hct: very high WBC, high concentration large PLT, autoagglutinins</td><td>MCV: very high WBC count, high concentration of very large PLTs, agglutinin RBCs, RBC fragments <36 fL, rigid RBCs</td><td>MCV and Hct: very high WBC, high concentration of large PLT, autoagglutinins</td></tr><tr><td>Platelet</td><td>very small RBCs and WBC fragments may interfere</td><td>very small red cells near the upper threshold, cell fragments, clumped PLTs, PLT fragments or cellular</td><td>very small RBCs and WBC fragments may cause no fit</td></tr><tr><td></td><td></td><td>debris near the lower PLT threshold, giant PLTs, PLT clumps, red and white cell fragments, electronic noise, verv small red cells</td><td></td></tr><tr><td>• Hemoglobin</td><td>very high WBC, severe lipemia, heparin, rare lyse-resistant RBCs</td><td>very high WBC count, severe lipemia, heparin, lyse-resistant RBCs, turbidity such as elevated trialycerides</td><td>very high WBC, severe lipemia, heparin, rare lyse-resistant RBCs</td></tr><tr><td>Interfering substances: Differential</td><td>high triglycerides may affect lysing</td><td>factors that affect WBC count above or high triglyc- erides that affect lysing, hypogranulas granulocytes, agranular granulocytes, lyse-resistant red cells, very</td><td>high triglycerides may affect lysing</td></tr><tr><td></td><td></td><td>small or multi-population lymphocytes, elevated triglycerides, precipitated elevated proteins</td><td></td></tr><tr><td>Maximum CBCs per hour/Maximum CBCs and differentials per hour</td><td>110/110</td><td>75/75</td><td>75/75</td></tr><tr><td>Minimum specimen volume open/Closed/Sample dead volume closed Microsample capability</td><td>200 μL/300 μL, 550 μL with slidemaker/1.0 mL ves</td><td>125 μL/185 μL/tube dependent ves</td><td>125 μL/185 μL/50 μL predilute/0.5 mL ves</td></tr><tr><td>Prepares microscope slides automatically or flags problems for slide prep</td><td>yes, both</td><td>no</td><td>no</td></tr><tr><td>Number of automatic slidemakers available/List price</td><td>>1,000 (U.S.)/\$110,000</td><td>_</td><td>-</td></tr><tr><td>Archives patient data/Previous patient results incl. with recent results</td><td>yes/yes</td><td>yes/yes</td><td>yes</td></tr><tr><td>Naximum archived data accessible when system online No. specimens for which numeric results saved in memory at once</td><td>20,000 samples 20,000 samples</td><td>20,000 samples 20,000 samples</td><td>5,000 samples 5,000 samples</td></tr><tr><td>No. specimens for which histo/cytogram results saved in memory at once Performs delta checks</td><td>20,000 samples ves</td><td>20,000 samples ves</td><td>5,000 samples</td></tr><tr><td>Tags and holds results for followup, confirmatory testing, or rerun</td><td>yes</td><td>yes</td><td>yes</td></tr><tr><td>Scattergram display: cell-specific color</td><td>yes</td><td>yes</td><td>four colors-cell types</td></tr><tr><td>Histogram display: color with thresholds User interface can display choice of specimen/result information</td><td>yes yes</td><td>yes yes</td><td>colors without thresholds no</td></tr><tr><td>LIS interface formats supported</td><td>RS-232. proprietary</td><td>RS-232, proprietary</td><td>RS-232, proprietary</td></tr><tr><td>Information transferred on LIS interface</td><td>numeric and flag results, histograms and</td><td>numeric and flag results, histograms and</td><td>numeric and flag results, histograms and</td></tr><tr><td></td><td>demographics, orders, LIS to instrument—</td><td>demographics, orders, LIS to instrument—</td><td>demographics, orders, LIS to instrument—</td></tr><tr><td>LOINC codes transmitted with all results/Sent in message to LIS/</td><td>broadcast no/no/no</td><td>broadcast no/no/no</td><td>broadcast no/no/no</td></tr><tr><td>Listing of machine codes and corresponding LOINC for each test</td><td>Bookman Coultor</td><td></td><td>_</td></tr><tr><td>Bar-code symbologies read on specimen tube</td><td>Codabar, codes 39 and 128, Interleaved 2 of 5, NW7</td><td>Codabar, codes 39 and 128, Interleaved 2 of 5, NW7</td><td>Codabar, codes 39 and 128, Interleaved 2 of 5, NW7</td></tr><tr><td>Accommodates bar-code placement per CLSI standard Auto2A</td><td>yes</td><td>yes</td><td>no</td></tr><tr><td>Time required for maintenance by lab personnel</td><td>no routine maintenance; only as needed</td><td>no routine maintenance; only as needed</td><td>none</td></tr><tr><td>Onboard diagnostics for troubleshooting/Limited to software problems Manufacturer can perform diagnostics via modem</td><td>yes/no yes</td><td>yes/no yes</td><td>yes/no no</td></tr><tr><td>Distinguishing features (supplied by company)</td><td>extensive decision support; enumeration of NRBCs with every differential: random access; automation</td><td>extensive decision support, extended linearity for WBC and PLT, low review rate, small footprint</td><td>VCS technology; low review rate; no routine daily maintenance; triplicate counting; aperture burn</td></tr><tr><td></td><td>ready; extended linearity for WBC and PLTs; RUO:</td><td>superior reliability, ProService, electronic IQAP</td><td>circuit; sweepflow; SmartStart system; autoloader</td></tr><tr><td>Note: a dash in lieu of an answer means company did not answer question or question is not applicable</td><td>יישט חרש</td><td></td><td>מווע אוועוכ־סמווµול וווטעלוא</td></tr></tbody></table>		

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Hematology analyzers			
Part 4 of 11	Beckman Coulter Bill Bailey bdbailey@beckman.com 250 South Kraemer Blvd.	CellaVision Ron Hagner ron.hagner@cellavision.com 4107 Burns Road	HORIBA Medical Jim Knowles jimknowles@horiba.com 34 Bunsen
See captodayonline.com/productguides for an interactive version of guide	Brea, CA 92821 714-961-4440 www.beckmancoulter.com	Beach Gardens, FL 33410 919-619-3909 cellavision.com	Irvine, CA 92618 888-903-5001 ext. 4553 www.horiba.com/us/en/medical
Name of instrument First year installed in U.S./Outside U.S./No. of units sold in 2011 No. units installed in U.S./Outside U.S./List price	Coulter Ac•T 5diff Family; Ac•T 5diff AL 2001/2000; 2003/2003/— >1,300/>2,800 combined in and out US \$38,500 (CP)/54,500 (AL)	CellaVision DM96 and CellaVision DM1200 2004/2003/— ~400/~700/~\$135,000-\$175,000	Pentra 60C+ Hematology Analyzer 2000/2000/85 >350/>600/\$47,313
Test menu: • Chartable (standard menu: WBC, RBC, Hb, Hct, MCV, MCH, MCHC, PLT, %&# neut, mono, lymph, eos, baso):	standard menu (left) plus: RDW, MPV	%&# neut, mono, lymph, eos, baso, segmented, bands, blast, promyelocytes, myelocytes, metamyelocytes, variant lymphocytes, plasma cells, giant platelets, platelet clumps, erythroblasts RBC morphology pre-characterizations include anisocytosis, poikilocytosis, polychromasia,</td><td>standard menu (left) plus: RDW, MPV</td></tr><tr><td>• Laboratory</td><td>atypical lymphocytes # (ATL#), atypical lymphocytes % (ATL%), immature cells # (IMM#), immature cells % (IMM%), PCT, PDW</td><td>microcytosis, macrocytosis, hypochromia —</td><td>atypical lymphocytes, atypical lymphocytes %, LIC, LIC %</td></tr><tr><td>• Flags</td><td>complete operator selectable flagging</td><td>_</td><td>operator selectable flagging</td></tr><tr><td>FDA-cleared tests not clinically released Tests not available but submitted for 510(k) clearance Tests in development Tests for research use only</td><td>— — PCT, PDW, IMM, ATL —</td><td>— — — analysis of cytocentrifuged samples body fluids</td><td> PCT, PDW, ATL, LIC </td></tr><tr><td></td><td></td><td>(Reported parameters: neutrophils, eosinophils, lymphocytes, macrophages (including monocytes), other (basophils, lymphoma cells, atypical lymphcytes, blast cells, and tumor cells)</td><td></td></tr><tr><td>Differential method(s) used</td><td>ACV technology combining cytochemistry, focused</td><td>light microscopy, image analysis and artificial neural networks</td><td>DHSS technology combining cytochemistry, focused flow impedance, and light absorbance principles of</td></tr><tr><td>Linearity: • WBC count/RBC count</td><td>measurement 0.4–91.3/.23-7.7; AL: 0.4–120.0/0.3–8.0</td><td></td><td>measurement 0–120/0–8</td></tr><tr><td>Hemoglobin/Platelet MCV (fL) or Hct (%) Precision: WBC count/RBC count Hemoglobin/Platelet MCV or Hct Accuracy of automated differential compared with manual differential (per CLSI H-20A2)</td><td>0–22/4-1000; AL: 1.3–24.0/10.0–1,000 1.8–63.8 (Hct)* <2 percent/<2 percent <1 percent/<5 percent <2.0 percent (Hct); AL: <2.0% (Hct) not available in NCCLS H-20A format</td><td></td><td>0-24/0-1,900 0-67 (Hct) <2 percent/<2 percent <1 percent/<5 percent <2 percent (Hct) neut% r=0.99,; lymph% r=0.98,; mono% r=0.96,; eos% r=0.89,; baso% r=0.54,</td></tr><tr><td>Interfering substances: • WRC</td><td>NDDCc DIT dumns large DITs luce_resistant DDCs</td><td>y=0.97+0.2, r=0.941, band neut% y=0.87x+0.1, r=0.917</td><td>NDDCo DIT olumno luco-resistant DDCo</td></tr><tr><td>• RBC</td><td>cold agglutinins, PLT clumps, WBC overlinearity</td><td>_</td><td>cold agglutinins</td></tr><tr><td>MCV or Hct Platelet Hemoglobin Interfering substances: Differential</td><td>Hct: lipemic samples, high WBC, cold agglutinins RBC and WBC fragments elevated WBC, lipemia lyse-resistant RBCs, NRBCs, lipemia</td><td></td><td>Hct: extreme leukocytosis microcytes, PLT clumps extreme lipemia/leukocytosis NRBC, lyse-resistant RBCs, extreme hyperbilirubinemia</td></tr><tr><td>Maximum CBCs per hour/Maximum CBCs and differentials per hour Minimum specimen volume open/Closed/Sample dead volume closed</td><td>60/60; 80/80 30 μL for CBC/30 μL/varies by tube size; 53 μL for CBC differential/53 μL for CBC differential/varies by tube size</td><td>—/35 differentials —</td><td>60/60 30 μL for CBC and 53 μL for CBC and differential/30 μL for CBC and 53 μL for CBC and differential/—</td></tr><tr><td>Microsample capability Prepares microscope slides automatically or flags problems for slide prep</td><td>yes no</td><td>=</td><td>yes yes</td></tr><tr><td>Number of automatic slidemakers available/List price</td><td>_</td><td></td><td>_</td></tr><tr><td>Archives patient data/Previous patient results incl. with recent results</td><td>yes</td><td>yes/no</td><td>yes/yes, with MultiLink Data Manager</td></tr><tr><td>Maximum archived data accessible when system online No. specimens for which numeric results saved in memory at once No. specimens for which histo/cytogram results saved in memory at once Performs delta checks</td><td>10,000 samples 10,000 samples 10,000 samples no</td><td>unlimited ~4,000 — no</td><td>100,000 unlimited with backup unlimited with backup yes</td></tr><tr><td>Tags and holds results for followup, confirmatory testing, or rerun Parameters for flags for holding samples defined by user or vendor Scattergram display: cell-specific color</td><td>yes user or vendor no</td><td></td><td>yes user ves</td></tr><tr><td>Histogram display: color with thresholds User interface can display choice of specimen/result information</td><td>yes yes</td><td></td><td>yes yes</td></tr><tr><td>LIS interface formats supported Information transferred on LIS interface</td><td>proprietary; proprietary ASTM numeric and flag results, histograms and differential plots, instrument to LIS; patient demographics, orders, LIS to instrument— broadcast</td><td>ASTM 1394 numeric and flag results, instrument to LIS; patient demographics, orders, LIS to instrument— broadcast; host query for patient demographics and orders (when bar code is read, host is queried</td><td>ASTM 1394 and 1238, HL7, IEEE MIB numeric and flag results, histograms and scatter- plots, instrument to LIS; patient demographics, LIS to instrument—broadcast</td></tr><tr><td>LOINC codes transmitted with all results/Sent in message to LIS/ Listing of machine codes and corresponding LOINC for each test Interface available or planned to automated specimen-handling system</td><td>no/no/no no</td><td>for orders no/no/no</td><td>yes/yes/yes no</td></tr><tr><td>Bar-code symbologies read on specimen tube</td><td>Codabar, codes 39 and 128, Interleaved 2 of 5, EAN 8 and 13 ves</td><td>Codabar, codes 39 and 128, Interleaved 2 of 5, QR, DataMatrix</td><td>Codabar, codes 39 and 128, ASTM, Interleaved 2 of 5</td></tr><tr><td>Time required for maintenance by lab personnel</td><td>none</td><td>daily: none; weekly: 5 minutes</td><td>weekly: 15 minutes</td></tr><tr><td>Onboard diagnostics for troubleshooting/Limited to software problems Manufacturer can perform diagnostics via modem</td><td>yes/no no</td><td>yes/no no</td><td>yes/yes yes, with Data Manager</td></tr><tr><td>Distinguishing features (supplied by company) Note: a dash in lieu of an answer means company did not answer question or question is not applicable</td><td>quantitative five-part WBC differential; aspirates only 30 μL of sample; requires small space footprint and runs quietly; AL has auto repeat based on decision rules</td><td>fully automated slide handling and oiling available in two models for medium and large laboratories; performs peripheral blood and body fluid differentials; WBC and other nucleated cells classified into 18 different categories; RBC morphology characterized for six categories; network use allows remote review of blood smears and linking of multiple analyzers in multiple locations</td><td>reliable five-part WBC differential technology— MTBF more than 200 days; small footprint; small sample size of 53 µL</td></tr></tbody></table>	

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Hematology analyzers			
Part 5 of 11	HORIBA Medical	HOBIBA Medical	Medica Corporation
	Jim Knowles jimknowles@horiba.com	Jim Knowles jimknowles@horiba.com	Ray Morrill rmorrill@medicacorp.com
	34 Bunsen Irvine, CA 92618	34 Bunsen Irvine, CA 92618	5 Oak Park Drive Bedford, MA 01730
See captodayonline.com/productguides	888-903-5001 ext. 4553	888-903-5001 ext. 4553	781-541-7413
for an interactive version of guide	www.horiba.com/us/en/medical	www.horiba.com/us/en/medical	www.medicacorp.com
Name of instrument	Pentra XL 80	Pentra DX120	EasyCell assistant
First year installed in U.S./Outside U.S./No. of units sold in 2011	2004/2003/31 >250/>900/\$76.808	2005/2004/6 >20/>400/\$207 560	2010/2012/— —/—/\$55.000
	~200/~000/#10,000	20/2400/0407,000	/ / 400,000
Test menu: • Chartable (standard menu: WBC, BBC, Hb Hct MCV	standard menu (left) plus: automatic dilution of	standard menu (left) plus: NRRCs, reticulocytes	WRC RRC Pit %&# neut mono lymnh eos haso
MCH, MCHC, PLT, %&# neut, mono, lymph, eos, baso):</td><td>overrange results (WBC \times 3, RBC/Hgb/PLT \times 2),</td><td>IRF, MRV</td><td>% & # of segmented neutrophils, band neutrophils,</td></tr><tr><td></td><td>RDW, MPV</td><td></td><td>lymphocytes, monocytes, eosinophils, basophils, variant lymphocytes, NRBCs, smudge cells, more</td></tr><tr><td></td><td></td><td></td><td></td></tr><tr><td></td><td></td><td></td><td></td></tr><tr><td>Laboratory</td><td>atypical lymphocytes, atypical lymphocytes%, LIC,</td><td>LIC%&#, atypical lymphocytes %&#, IMG %&#, IML</td><td>-</td></tr><tr><td></td><td></td><td>MRU, MFI%, CRC%</td><td></td></tr><tr><td>• Flags</td><td>operator selectable flagging</td><td>—</td><td>-</td></tr><tr><td></td><td></td><td></td><td></td></tr><tr><td>FDA-cleared tests not clinically released Tests not available but submitted for 510(k) clearance</td><td></td><td></td><td>_</td></tr><tr><td>Tests in development</td><td><u> </u></td><td></td><td>_</td></tr><tr><td>Tests for research use only</td><td>PCT, PDW, ATL, LIC</td><td>PCT, PDW, ATL, LIC, IMG, IML, IMM</td><td>-</td></tr><tr><td>Tests unique to analyzer</td><td>automatic dilution protocol</td><td>-</td><td>competency assessment program</td></tr><tr><td></td><td></td><td></td><td></td></tr><tr><td></td><td></td><td></td><td></td></tr><tr><td></td><td></td><td></td><td></td></tr><tr><td>Differential method(s) used</td><td>DHSS technology combining cytochemistry, focused</td><td>cytochemistry (chlorazol black E) and absorbance</td><td>light microscopy, image analysis by neural network</td></tr><tr><td></td><td>flow impedance, and light absorbance</td><td></td><td>software, images reviewed by technologist</td></tr><tr><td>Linearity: • WBC count/RBC count</td><td>0–120/0–8</td><td>0–150/0.5–8.1</td><td>-</td></tr><tr><td>Hemoglobin/Platelet</td><td>0–24/0–1,900 (>2 g/dL Hb)</td><td>2–25/0–2,000</td><td>_</td></tr><tr><td>MCV (fL) or Hct (%) WPC count</td><td>0-67 (Hct)/0-2,800 (<2 g/dL Hb)</td><td>0-80 (Hct)</td><td>-</td></tr><tr><td>Hemoglobin/Platelet</td><td><2 percent/<2 percent <1 percent/<5 percent</td><td><2 percent/<2 percent <1 percent/<5 percent</td><td></td></tr><tr><td>MCV or Hct Accuracy of automated differential compared with manual</td><td><2 percent (Hct)</td><td><2 percent (Hct) neut% r=0.00 lymph% r=0.08 mono%</td><td></td></tr><tr><td>differential (per CLSI H-20A2)</td><td>r=0.96, —; eos% r=0.89, —; baso% r=0.54, —</td><td>r=0.92, -; eos% r=0.97, -; baso% r=0.71, -</td><td>r=0.93; eosinophil r=0.97</td></tr><tr><td></td><td></td><td></td><td></td></tr><tr><td>Interfering substances: • WBC</td><td>NRBCs, PLT clumps, lyse-resistant RBCs</td><td>NRBCs, PLT clumps, lyse-resistant RBCs</td><td>-</td></tr><tr><td>• BBC</td><td>cold applutining</td><td>cold applutining</td><td>_</td></tr><tr><td>MCV or Hct</td><td>Hct: extreme leukocytosis</td><td>Hct: extreme leukocytosis</td><td>-</td></tr><tr><td> Platelet Hemoalobin </td><td>microcytes, PLT clumps extreme lipemia. leukocytosis</td><td>microcytes, PLT clumps extreme lipemia. leukocytosis</td><td><u> </u></td></tr><tr><td>Interfering substances: Differential</td><td>NRBCs, lyse-resistant RBCs, extreme</td><td>NRBCs, lyse-resistant RBCs, extreme</td><td>_</td></tr><tr><td></td><td></td><td>nyperbilirubinemia</td><td></td></tr><tr><td>Maximum CBCs per hour/Maximum CBCs and differentials per hour Minimum specimen volume open/Closed/Sample dead volume closed</td><td>80/80 30 ul for CBC/53 ul for CBC and differential/0.5 ml</td><td>120/120 130 ul /200 ul /1 ml</td><td>—/12 —</td></tr><tr><td></td><td></td><td></td><td></td></tr><tr><td>Microsample capability Prepares microscope slides automatically or flags problems for slide prep</td><td>yes ves</td><td>yes, open mode ves</td><td> no</td></tr><tr><td></td><td>,</td><td>,</td><td></td></tr><tr><td>Number of automatic slidemakers available/List price</td><td></td><td></td><td></td></tr><tr><td>Archives patient data/Previous patient results incl. with recent results</td><td>yes/yes, with MultiLink Data Manager</td><td>yes/yes, with MultiLink Data Manager</td><td>yes/no</td></tr><tr><td>Maximum archived data accessible when system online No. specimens for which numeric results saved in memory at once</td><td>100,000 unlimited with backup</td><td>100,000 unlimited with backup</td><td>unlimited 10.000</td></tr><tr><td>No. specimens for which histo/cytogram results saved in memory at once</td><td>unlimited with backup</td><td>unlimited with backup</td><td></td></tr><tr><td>Tags and holds results for followup, confirmatory testing, or rerun</td><td>yes yes</td><td>yes yes</td><td>no </td></tr><tr><td>Parameters for flags for holding samples defined by user or vendor</td><td>user</td><td>user</td><td>-</td></tr><tr><td>Histogram display: color with thresholds</td><td>yes yes</td><td>yes yes</td><td>_</td></tr><tr><td>User interface can display choice of specimen/result information</td><td>-</td><td>yes</td><td>_</td></tr><tr><td>LIS interface formats supported</td><td>proprietary, ASTM 1394 and 1238, HL7, IEEE MIB</td><td>proprietary, ASTM 1394 and 1238, HL7, IEEE MIB</td><td>ASTM 1394, LAN connection allows remote review</td></tr><tr><td>Information transformed on LIS interface</td><td>numorio and flag results, histograms and seat-</td><td>numeric and flag results, histograms and seat-</td><td>of all slide images</td></tr><tr><td></td><td>terplots, instrument to LIS; patient demographics,</td><td>terplots, instrument to LIS; patient demographics,</td><td>patient demographics, orders, LIS to instrument—</td></tr><tr><td></td><td>orders, LIS to instrument— broadcast</td><td>orders, LIS to instrument— broadcast</td><td>broadcast; host query for demographics and orders</td></tr><tr><td></td><td></td><td></td><td></td></tr><tr><td>LUINC codes transmitted with all results/Sent in message to LIS/ Listing of machine codes and corresponding LOINC for each test</td><td>yes/yes/yes</td><td>no/no/no</td><td>yes/yes/yes</td></tr><tr><td>Interface available or planned to automated specimen-handling system</td><td>yes</td><td>yes</td><td><u>-</u></td></tr><tr><td>Bar-code symbologies read on specimen tube</td><td>Codabar, codes 39 and 128, ASTM, Interleaved 2 of 5</td><td>Codabar, codes 39 and 128, ASTM, Interleaved 2 of 5</td><td>Codabar, codes 39 and 128, Interleaved 2 of 5, PDF- 417 (two dimensional)</td></tr><tr><td>Accommodates bar-code placement per CLSI standard Auto2A</td><td>yes</td><td>yes</td><td></td></tr><tr><td>Time required for maintenance by lab personnel</td><td>weekly: 15 minutes</td><td>weekly: 15 minutes</td><td>daily: 5 minutes; weekly: 10 minutes</td></tr><tr><td>Onhoard diagnostics for troublochooting // imited to optimize methods</td><td>no/ves</td><td>no/ves</td><td>ves/no</td></tr><tr><td>Manufacturer can perform diagnostics via modem</td><td>yes</td><td>yes</td><td>no</td></tr><tr><td>Distinguishing features (supplied by company)</td><td>compact five-part differential instrument with</td><td>high-throughout call counter with integrated</td><td>30-nosition slide autosampler for walkowov opera</td></tr><tr><td>sounguonny routers (supplied by company)</td><td>autoloader and autodilution capability, auto rerun</td><td>reticulocyte methodology and slidemaker-stainer;</td><td>tion and separate stat position to allow immediate</td></tr><tr><td></td><td>feature, autovalidation</td><td>fluorescent NRBC counting, auto rerun and reflex testing, autovalidation</td><td>analysis of stat sample; remote review software creates additional workstations, which improves</td></tr><tr><td></td><td></td><td></td><td>workflow efficiency: allows easy collaboration on</td></tr><tr><td></td><td></td><td></td><td>difficult clides from clining to the second clides from clining</td></tr><tr><td></td><td></td><td></td><td>difficult slides from alternate locations outside the laboratory; slides without bar codes are imaged for</td></tr></tbody></table>			

Note: a dash in lieu of an answer means company did not answer question or question is not applicable

Hematology analyzers

Part 6 of 11	Siemens Healthcare Diagnostics Lois Brisben lois.brisben@siemens.com 1717 Deerfield Road Deerfield II 60015-0778	Siemens Healthcare Diagnostics Lois Brisben lois.brisben@siemens.com 1717 Deerfield Road Deerfield II 60015-0778	Siemens Healthcare Diagnostics Lois Brisben lois.brisben@siemens.com 1717 Deerfield Road Deerfield III 60015-0778
See captodayonline.com/productguides for an interactive version of guide	800-948-3234 www.usa.siemens.com/diagnostics	800-948-3234 www.usa.siemens.com/diagnostics	800-948-3234 www.usa.siemens.com/diagnostics
Name of instrument First year installed in U.S./Outside U.S./No. of units sold in 2011 No. units installed in U.S./Outside U.S./List price	Advia 120 Hematology System 1998/1998/— >750/3,500/\$169,000–\$189,000	Advia 2120 Hematology System 2004/2004/— >200/>900/\$225,000	Advia 2120i 2008/2008/130 >150/>400/\$225,000
Test menu: • Chartable (standard menu: WBC, RBC, Hb, Hct, MCV, MCH, MCHC, PLT, %&# neut, mono, lymph, eos, baso): • Laboratory • Flags	standard menu (left) plus: CHCM, MPV, RDW, HDW, LUC %&#, retic %&#, CHr, CHCMr, MCVr; CSF: WBC, RBC, PMN, MN, neut, lymph, mono; cellular Hgb %: hypo, hyper, macro, micro; calc. Hb, MPXI; %: blasts, PMN, MN; large PLT count; RBC fragment count; RBC ghost count; CSF: WBC, RBC, three-part differential; body fluids: TNC, RBC left shift, atyp. lymph, blasts, immature grans, myeloperoxidase deficiency, aniso, micro, macro, Hb variation, hypo, hyper, NRBC, RBC fragments, RBC ghost, large PLT, PLT clumps</td><td>standard menu (left) plus: CHCM, MPV, RDW, HDW, LUC %&#, retic %&#, CHr, CHCMr, cellular Hgb, MCVr; CSF: WBC, RBC, PMN, MN, neut, lymph, mono % hypo, hyper, macro, micro; MPXI, %: blast, PMN, MN, large PLT count, RBC fragment count; RBC ghost count; NRBC; CSF: WBC, RBC, three-part differential; body fluids: TNC, RBC left shift, atyp. lymph, blasts, immature grans, myeloperoxidase deficiency, aniso, micro, macro, Hb variation, hypo, hyper, NRBC, RBC fragments, RBC ghost, large PLT, PLT clumps</td><td>standard menu (left) plus: CHCM, MPV, RDW, HDW, LUC %&#, retic. %&#, CHr, CHCMr, cellular Hgb, MCVr; CSF: WBC, RBC, PMN, MN, neut, lymph, mono %hypo, hyper, macro, micro, MPXI, %blast, PMN, MN, large PLT count, RBC fragment count, RBC ghost count, NRBC; CSF: WBC, RBC, three-part differential; body fluids: TNC, RBC left shift, atypical lymphocytes, blasts, immature grans, myeloperoxidase deficiency, aniso, micro, macro, Hgb variation, hypo, hyper, NRBC, RBC fragments, RBC ghost, large PLT, PLT clumps</td></tr><tr><td>FDA-cleared tests not clinically released Tests not available but submitted for 510(k) clearance Tests in development Tests for research use only Tests unique to analyzer</td><td>— — IRF, MPC, MPM CSF, eos CHCM, HDW, CHr, CHCMr, MPC, MPM; CSF: WBC RBC, MN, PMN, neut, lymph, mono</td><td>— — MPC, MPM IRF, CSF, eos CHCM, HDW, CHr, CHCMr, cellular Hgb, MPC, MPM, CSF: WBC, RBC, PMN, MN, neut, lymph, mono</td><td>— — MPC, MPM IRF, CSF eos CHCM, HDW, CHr, CHCMr, cellular Hgb, MPC, MPM, CSF: WBC, RBC, PMN, MN, neut, lymph, mono</td></tr><tr><td>Differential method(s) used Linearity: • WBC count/RBC count • Hemoglobin/Platelet • MCV (fL) or Hct (%) Precision: • WBC count/RBC count • Hemoglobin/Platelet • MCV or Hct Accuracy of automated differential compared with manual differential (per CLSI H-20A2) Interfering substances: • WBC</td><td>perox-peroxidase cytochemistry staining with light scatter and absorption; baso-cytochemistry stripping with two-angle laser light scatter $0.02-400/0-7.0$; CSF WBC $0-5,000/\mu$L; CSF RBC $0-1,500/\mu$L 0-22.5/5-3,500 30-180</math> (MCV) 2.7 percent/1.2 percent 0.93</math> percent/2.93 percent 0.78</math> percent (MCV) neut% r=0.997, y=1.02x-0.6; lymph% r=0.997, y=1.00x+0.8; mono% r=0.943, y=0.85x-0.3; eos% r=0.979, y=0.87x+0.2; baso% r=0.772, y=0.67x+0.0; luc% r=0.994, y=0.92x+0.6 incomplete RBC lysis (peroxidase only)</td><td>peroxidase WBC—peroxidase cytochem. staining with light scatter and absorption; baso—cytochem. stripping with two-angle laser light scatter 0.02-400</math>; CSF WBC 0-5,000/0-7.0; CSF RBC 0-15,00 0-22.5/5-3,500 30-180</math> (MCV) 2.7 percent/1.2 percent 0.93</math> percent/2.93 percent 0.78</math> percent (MCV) neut% r=0.997, y=1.02x-0.6; lymph% r=0.997, y=1.00x+0.8; mon0% r=0.943, y=0.85x-0.3; eos% r=0.979, y=0.87x+0.2; baso% r=0.772, y=0.67x+0.0; luc% r=0.994, y=0.92x+0.6 incomplete RBC lysis (peroxidase only)</td><td>peroxidase WBC: peroxidase cytochem. staining with light scatter and absorption; baso: cytochem. stripping with two-angle laser light scatter 0.02-400 CSF: 0-5,000/0-7.0 CSF: 0-1,500 0-22.5/5-3,500 30-180 (MCV) 2.7 percent/1.2 percent 0.93 percent/2.93 percent 0.78 percent (MCV) neut% r=0.997, y=1.02x-0.6; lymph% r=0.997, y=1.00x+0.8; mon0% r=0.943, y=0.85x-0.3; eos% r=0.979, y=0.87x+0.2; baso% r=0.772, y=0.67x+0.0; luc% r=0.994, y=0.92+0.6 incomplete RBC lysis (peroxidase only)</td></tr><tr><td>• RBC</td><td>cold agglutinins, extreme sickle cell</td><td>cold agglutinins, extreme sickle cell</td><td>cold agglutinins, extreme sickle cell</td></tr><tr><td>MCV or Hct</td><td>none</td><td>_</td><td>none</td></tr><tr><td>Platelet</td><td>none</td><td>_</td><td>none</td></tr><tr><td>Hemoglobin Interfering substances: Differential</td><td>high WBC, lipemin, extremely high bilirubin, interfere with cyanmethemoglobin only, none with direct cellular Hb (CHCM) incomplete lysis of RBCs, complete myeloperoxidase deficiency</td><td>extreme lipemia, high WBC, extremely high bilirubin interference with colorimetric Hb only, none with cellular Hb incomplete RBC lysis, complete myeloperoxidase deficiency</td><td>extreme lipemia, high WBC, extremely high bilirubin—interference with colorimetric Hgb only, none with cellular Hgb incomplete RBC lysis, complete myeloperoxidase deficiency</td></tr><tr><td>Maximum CBCs per hour/Maximum CBCs and differentials per hour Minimum specimen volume open/Closed/Sample dead volume closed</td><td>120/120 157 μL/157 μL/<300 μL (tube size dependent)</td><td>120/120 175 μL/175 μL/<300 (tube size dependent)</td><td>120/120 175 μL/175 μL/<300 (tube size dependent)</td></tr><tr><td>Microsample capability Prepares microscope slides automatically or flags problems for slide prep</td><td>yes yes</td><td>yes if integrated to Advia Autoslide</td><td>yes yes</td></tr><tr><td>Number of automatic slidemakers available/List price</td><td></td><td>Advia Autoslide, —/\$98,000</td><td>Advia Autoslide, —/\$98,000</td></tr><tr><td>Archives patient data/Previous patient results incl. with recent results Maximum archived data accessible when system online No. specimens for which numeric results saved in memory at once No. specimens for which histo/cytogram results saved in memory at once Performs delta checks Tags and holds results for followup, confirmatory testing, or rerun Parameters for flags for holding samples defined by user or vendor Scattergram display: cell-specific color Histogram display: color with thresholds User interface can display choice of specimen/result information</td><td>yes/no 10,000 samples 10,000 samples 10,000 samples yes yes user or vendor yes yes yes</td><td>yes/no 10,000 10,000 yes yes user or vendor yes yes yes yes</td><td>yes/no 10,000 samples 10,000 samples 10,000 samples yes yes yes yes yes yes yes</td></tr><tr><td>LIS interface formats supported Information transferred on LIS interface</td><td>proprietary (Spec 79) numeric and flag results, histograms and scatterplots, instrument to LIS; patient demographics, orders, LIS to instrument— broadcast; host query for demographics and orders</td><td>proprietary numeric and flag results, histograms and scatterplots, instrument to LIS; patient demographics, patient orders, LIS to instrument— broadcast; host query for patient demographics and orders (when bar code is read, host is queried for orders)</td><td>proprietary (instrument or vendor specific) numeric and flag results, histograms and scatterplots, instrument to LIS; patient demographics, orders, LIS to instrument— broadcast; host query for demographics and orders</td></tr><tr><td>LOINC codes transmitted with all results/Sent in message to LIS/ Listing of machine codes and corresponding LOINC for each test Interface available or planned to automated specimen-handling system Bar-code symbologies read on specimen tube</td><td>no/no/yes LabCell (Siemens) Codabar, codes 39 and 128, ASTM, Interleaved 2 of 5</td><td>no/no/yes LabCell (Siemens) Codabar, codes 39 and 128, Interleaved 2 of 5</td><td>no/no/yes LabCell (Siemens) Codabar, codes 39 and 128, ASTM, Interleaved 2 of 5</td></tr><tr><td>Accommodates bar-code placement per CLSI standard Auto2A</td><td>yes</td><td></td><td>yes</td></tr><tr><td>Time required for maintenance by lab personnel</td><td>daily: 10 minutes; weekly: 15 minutes; monthly: 15 minutes</td><td>weekly: 15 minutes; monthly: 15 minutes</td><td>daily: 10 minutes; weekly: 15 minutes; monthly: 15 minutes</td></tr><tr><td>Onboard diagnostics for troubleshooting/Limited to software problems Manufacturer can perform diagnostics via modem</td><td>yes/no yes</td><td>yes/no yes</td><td>yes/no yes</td></tr><tr><td>Distinguishing features (supplied by company)</td><td>laser technology provides cellular Hb for RBCs and retics; 2-D PLT analysis eliminates interference from RBC fragments and inclusion of large PLTs; dual WBC counts with a linearity of up to 400,000; CSF assay</td><td>laser technology provides direct cellular Hb for RBCs and reticulocytes; 2-D PLT analysis eliminates interference from RBC fragments and inclusion of large PLTs; dual WBC counts with a linearity of up to 400,000; CSF assay</td><td>laser technology provides direct cellular Hgb for RBCs and reticulocytes; 2-D PLT analysis eliminates interference from RBC fragments and inclusion of large PLTs; dual WBC counts with a linearity of up to 400,000; CSF assay</td></tr></tbody></table>		

Note: a dash in lieu of an answer means company did not answer question

December 2012

Hematology analyzers			
Part 7 of 11	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic Road	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic Road	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic Road
See captodayonline.com/productguides for an interactive version of guide	Lincolnshire, IL 60069 800-379-7639 www.sysmex.com/us	Lincolnshire, IL 60069 800-379-7639 www.sysmex.com/us	Lincolnshire, IL 60069 800-379-7639 www.sysmex.com/us
Name of instrument	Sysmex pocH-100i	Sysmex KX-21N	XS-1000i and XS-1000i AutoLoader (20-sample autoloader option)
First year installed in U.S./Outside U.S./No. of units sold in 2011 No. units installed in U.S./Outside U.S./List price	2004/2003/>130 >1,100/>4,400/\$19,094	2001/1999/125 >1,200/>4,800/\$28,408	2006/2005/>340 >1,800/>7,000/\$91,052 (XS-1000i) \$101,764 (AutoLoader)
Test menu: • Chartable (standard menu: WBC, RBC, Hb, Hct, MCV, MCH, MCHC, PLT, %&# neut, mono, lymph, eos, baso):	WBC, RBCs, Hb, Hct, MCV, MCH, MCHC, PLT, %&# neut, lymph, MXD (mono, eos, baso), RDW-SD, RDW-</td><td>WBC, RBC, Hb, Hct, MCV, MCH, MCHC, PLT, %&# neut, lymph, MXD (mono, eos, baso), RDW-SD, RDW-CV,</td><td>standard menu (left) plus: MPV, RDW-SD, RDW-CV</td></tr><tr><td>Laboratory</td><td>CV, MPV </td><td>MPV </td><td>-</td></tr><tr><td>• Flags</td><td>histogram error flags; WBC, RBC, PLT</td><td>histogram error flags; WBC, RBC, PLT</td><td>PLT clumps, PLT ABN distribution, WBC ABN scattergram, blast immature granulocytes, left shift, atypical lymphocytes, abnormal lymphocytes/ blasts, RBC ABN distribution, RBC lyse resistance, RBC agglutinins, turbidity, NRBC</td></tr><tr><td>FDA-cleared tests not clinically released Tests not available but submitted for 510(k) clearance</td><td>=</td><td>-</td><td></td></tr><tr><td>Tests in development Tests for research use only</td><td>— — </td><td>— — </td><td>IG% research screen</td></tr><tr><td>lests unique to analyzer</td><td>absolute neutrophil count</td><td>absolute neutrophil count</td><td>_</td></tr><tr><td>Differential method(s) used</td><td>direct current</td><td>direct current</td><td>fluorescent flow cytometry</td></tr><tr><td>Linearity: • WBC count/RBC count • Hemoglobin/Platelet</td><td>1.0–99.9/0.3–7.0 0.1–25.0/10–999</td><td>1.0–99.9/0.3–7.0 0.1–25.0/10–999</td><td>0–400/0–8 0-25/0–5,000</td></tr><tr><td>MCV (fL) or Hct (%) Precision: WBC count/RBC count</td><td>10–60 Hct <=3.5 percent/<=2.0 percent</td><td>10–60 Hct <=3.5 percent/<=2.0 percent</td><td>0–60 (Hct) —</td></tr><tr><td> Hemoglobin/Platelet MCV or Hct </td><td><=1.5 percent/<=6.0 percent <=2.0 percent Hct</td><td><=1.5 percent/<=6.0 percent <=2.0 percent Hct</td><td>_</td></tr><tr><td>Accuracy of automated differential compared with manual differential (per CLSI H-20A2)</td><td>NEUT% R=0.98, LYM% R=0.99, MXD % R=0.75, NEUT# R=1.00, LYM# R=1.00, MXD# R=0.90</td><td>NEUT% R=0.98, LYM% R=0.99, MXD % R=0.75, NEUT# R=1.00, LYM# R=1.00, MXD# R=0.90</td><td>neut% r=0.96, y=0.9074x+3.8948; lymph% r=0.97, y=0.9017x+2.4817; mono% r=0.78, y=0.8626x+3.5938; eos% r=0.94, y=0.9076x+0.3651; baso% r=0.29, y-0.1538x+0.298</td></tr><tr><td>Interfering substances: • WBC</td><td>lyse-resistant RBCs, cold agglutinins,</td><td>cold agglutinins, PLT aggregation, erythroblastosis,</td><td>cold agglutinins, PLT aggregation, cryoglobulins,</td></tr><tr><td>• RBC</td><td>cold agglutinins, rLi aggregation, NRBCS cold agglutinins, microcytosis (severe), fragmented RBCs</td><td>cold agglutinin, severe microcytosis, fragmented RBCs, leukocytosis (>100,000/µL)</td><td>cold agglutinins, severe microcytosis, fragmented RBCs, leukocytosis</td></tr><tr><td>• MCV or Hct</td><td>cold agglutinins, fragmented RBCs, leukocytosis (>100,000/uL)</td><td>Hct: cold agglutinin, leukocytosis (>100,000/µL), abnormal red cell fragility, spherocytosis</td><td>Hct: cold agglutinins, ABN red cell fragility, spherocytosis, leukocytosis (>100,000/µL)</td></tr><tr><td>Platelet</td><td>PLT aggregation, giant PLTs, microcytic RBCs, fragmented RBCs</td><td>pseudothrombocytopenia, PLT aggregation, increased microcytosis, megalocytic PLTs</td><td>pseudothrombocytopenia, PLT aggregation, increased microcytosis, megaloblasts</td></tr><tr><td>• Hemoglobin</td><td>lipemia (severe), abnormal protein, leukocytosis (>100,000/uL)</td><td>leukocytosis (>100,000/µL), lipemia, abnormal proteins</td><td>lipemia, ABN proteins, leukocytosis (>100,000/µL)</td></tr><tr><td>Interfering substances: Differential</td><td>_</td><td>_</td><td>lyse-resistant RBCs</td></tr><tr><td>Maximum CBCs per hour/Maximum CBCs and differentials per hour Minimum specimen volume open/Closed/Sample dead volume closed</td><td>30/30 15 μL/15 μL/15 μL</td><td>60/60 50 μL/—/—</td><td>60/60 20 μL/20 μL/1.0 mL</td></tr><tr><td>Microsample capability Prepares microscope slides automatically or flags problems for slide prep</td><td>yes no</td><td>yes no</td><td>yes no</td></tr><tr><td>Number of automatic slidemakers available/List price</td><td>_</td><td>_</td><td>_</td></tr><tr><td>Archives patient data/Previous patient results incl. with recent results Maximum archived data accessible when system online</td><td>yes/yes 100 samples</td><td>yes/yes 300 samples</td><td>yes/yes 10.000 samples</td></tr><tr><td>No. specimens for which numeric results saved in memory at once No. specimens for which histo/cytogram results saved in memory at once</td><td>100 samples 100 samples</td><td>300 samples 300 samples</td><td>10,000 samples 10,000 samples</td></tr><tr><td>Performs delta checks Tags and holds results for followup, confirmatory testing, or rerun</td><td>yes no</td><td>yes</td><td>yes</td></tr><tr><td>Parameters for flags for holding samples defined by user or vendor Scattergram display: cell-specific color</td><td>yes no</td><td>yes no</td><td>user or vendor ves</td></tr><tr><td>Histogram display: color with thresholds User interface can display choice of specimen/result information</td><td>yes ves</td><td>yes ves</td><td>yes ves</td></tr><tr><td>LIS interface formats supported Information transferred on LIS interface</td><td>RS-232C numeric and flag results, histograms and scatterplots, patient demographics, patient orders, host query for patient demographics and orders</td><td>RS-232C numeric and flag results, histograms and scatterplots, host query for patient demographics and orders</td><td>proprietary, ASTM 1394, TCP-IP numeric and flag results, histograms and scatterplots, patient demographics, orders</td></tr><tr><td>LOINC codes transmitted with all results/Sent in message to LIS/ Listing of machine codes and corresponding LOINC for each test</td><td>no/no/yes</td><td>no/no/yes</td><td>no/no/yes</td></tr><tr><td>Interface available or planned to automated specimen-handling system Bar-code symbologies read on specimen tube</td><td> codes 39 and 128, ASTM, ITF, NW7, JAN-8 and 13</td><td> codes 39 and 128, ITF, NW-7, JAN, UPC-A, UPC-E, EAN13. EAN8</td><td> Codabar, codes 39 and 128, ASTM, Interleaved 2 of 5. NW7, EAN 8 and 13. ITE</td></tr><tr><td>Accommodates bar-code placement per CLSI standard Auto2A</td><td>yes</td><td>yes</td><td>yes</td></tr><tr><td>Time required for maintenance by lab personnel</td><td>daily: <2 minutes; weekly: <2 minutes; monthly: <2 minutes</td><td>daily: <2 minutes; weekly: <2 minutes; monthly: <2 minutes</td><td>daily: 3 minutes; monthly: 9 minutes</td></tr><tr><td>Onboard diagnostics for troubleshooting/Limited to software problems Manufacturer can perform diagnostics via modem</td><td>yes/no yes</td><td>yes/no yes</td><td>yes/no yes, also via Internet</td></tr><tr><td>Distinguishing features (supplied by company)</td><td>hydrodynamic focusing, automatic floating discriminators, ISBT-compliant, data masking software for blood donor centers</td><td>automatic floating discriminators</td><td>standardized technology, reagents, controls, and operations to other X series analyzers; small sample volume requirements for CBC and five-part differential; remote diagnostics, online QC, discrete analysis, reagent monitoring, chartable report; remote calibration verification</td></tr></tbody></table>		

Tabulation does not represent an endorsement by the College of American Pathologists.

December 2012

Hematology analyzers			
Part 8 of 11 See captodayonline.com/productguides for an interactive version of guide	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic Road Lincolnshire, IL 60069 800-379-7639 www.sysmex.com/us	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic Road Lincolnshire, IL 60069 800-379-7639 www.sysmex.com/us	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic Road Lincolnshire, IL 60069 800-379-7639 www.sysmex.com/us
Name of instrument	Sysmex XT-1800i	Sysmex XT-2000i	Sysmex XT-4000i
First year installed in U.S./Outside U.S./No. of units sold in 2011 No. units installed in U.S./Outside U.S./List price	2002/2001/>70 >420/4,600/\$137,917	2002/2001/>70 >980/>5,200/\$158,430	2010/2009/>120 >250/>900/\$195,700
Test menu: • Chartable (standard menu: WBC, RBC, Hb, Hct, MCV, MCH, MCHC, PLT, %&# neut, mono, lymph, eos, baso):	standard menu (left) plus: MPV, RDW-SD, RDW-CV, immature granulocytes %&#</td><td>standard menu (left) plus: retic %&#, IRF, PLT-O, MPV, RDW-SD, RDW-CV, reticulocyte hemoglobin, immature granulocytes %&#</td><td>standard menu (left) plus: IG% and #, retic % and #, IRF, RET-He, PLT-O, BF: RBC/WBC/TC/two-part differential</td></tr><tr><td> Laboratory Flags FD∆-cleared tests not clinically released </td><td>— PLT clumps, PLT ABN distribution, WBC ABN scattergram, blast immature granulocytes, left shift, atypical lymphocytes, abnormal lymphocytes/ blasts, RBC ABN distribution, RBC lyse resistance, RBC agglutinins, turbidity, NRBC, body fluids —</td><td>— PLT clumps, PLT ABN distribution, WBC ABN scattergram, blast immature granulocytes, left shift, atypical lymphocytes, abnormal lymphocytes/ blasts, RBC ABN distribution, RBC lyse resistance, RBC agglutinins, turbidity, NRBC, body fluids —</td><td>— PLT clumps, PLT abnormal distribution, blast, imm grans, left shift, atypical lymphocytes, abnormal lymphocytes/blasts, NRBC, RBC lyse resistance, RBC ABN distribution, RBC agglutinins, turbidity —</td></tr><tr><td>Tests not available but submitted for 510(k) clearance Tests in development</td><td>_</td><td>_</td><td>-</td></tr><tr><td>Tests for research use only Tests unique to analyzer</td><td> immature granulocytes (IG%&#)</td><td> PLT-O, immature granulocytes (IG) %&#, reticulocyte hemoglobin (RET-He)</td><td> reticulocyte hemoglobin, immature reticulocyte fraction, reportable immature granulocyte # and %, PLT-O, BF: RBC, WBC, TC, two-part differential</td></tr><tr><td>Differential method(s) used Linearity: • WBC count/RBC count • Hemoglobin/Platelet • MCV (fL) or Hct (%) Precision: • WBC count/RBC count • Hemoglobin/Platelet • MCV or Hct Accuracy of automated differential compared with manual differential (per CLSI H-20A2)</td><td>fluorescent flow cytometry 0–310/0–8 0–25/0–5,000 0–60 (Hct) ≤3.0 percent/≤1.5 percent ≤1.5 percent/≤4.0 percent ≤1.5 percent(Hct) neut% r=0.95, y=0.95x+3.38; lymph% r=0.96, y=0.85x+1.67; mon0% r=0.90, y=11.37x+1.89; eos% r=0.94, y=0.87x+0.04; baso% r=0.76, y=0.48x+0.24</td><td>fluorescent flow cytometry 0-310/0-8 0-25/0-5,000 0-60</math> (Hct) $\leq 3.0 \text{ percent} \leq 1.5 \text{ percent}$ $\leq 1.5 \text{ percent} \leq 4.0 \text{ percent}$ $\leq 1.5 \text{ percent} (\text{Hct})$ neut% r=0.95, y=0.95x+3.38; lymph% r=0.96, y=0.85x+1.67; mono% r=0.90, y=11.37x+1.89; eos% r=0.94, y=0.87x+0.04; baso% r=0.76, y=0.48x+0.24</td><td>fluorescent flow cytometry 0-440/0-8 0-25/0-5,000 0-60 (Hct) <=3.0 percent/<=1.5 percent <=1.5 percent/<=4.0 percent <=1.5 percent (Hct) neut % r=0.95, lymph% r=0.96, mono% r=0.90, eos% r=0.94, baso% r=0.76; neut % y=0.95x+3.38, lymph % y=0.85x+1.67, mono % y=11.37x+1.89, eos% y=0.87x+0.04, baso% y=0.48x+0.24</td></tr><tr><td>Interfering substances: • WBC</td><td>cold agglutinins, PLT aggregation, cryoglobulins, lyse-resistant RBCs, NRBCs</td><td>cold agglutinins, PLT aggregation, cryoglobulins, lyse-resistant RBCs, NRBCs</td><td>cold agglutinins, PLT aggregation, cryoglobulins, lyse resistant erythrocytes, NRBCs</td></tr><tr><td>• RBC</td><td>cold agglutinins, severe microcytosis, fragmented RBCs, leukocytosis</td><td>cold agglutinins, severe microcytosis, fragmented RBCs, leukocytosis</td><td>cold agglutinins, severe microcytosis, fragmented RBCs, leukocytosis</td></tr><tr><td>• MCV or Hct</td><td>Hct: cold agglutinins, ABN red cell fragility, spherocytosis, leukocytosis (>100,000/µL)</td><td>Hct: cold agglutinins, ABN red cell fragility, spherocytosis. leukocytosis (>100,000/µL)</td><td>Hct: cold agglutinins, fragmented RBCs, spherocytosis, leukocytosis (lymphocytes>100,000/</td></tr><tr><td>• Platelet</td><td>pseudothrombocytopenia, PLT aggregations,</td><td>pseudothrombocytopenia, PLT aggregation,</td><td>μL) PLT aggregation, pseudothrombocytopenia, giant</td></tr><tr><td>• Hemoglobin</td><td>increased microcytosis, megaioplasts lipemia, ABN proteins, leukocytosis (>100,000/µL)</td><td>increased microcytosis, megaiobiasts lipemia, ABN proteins, leukocytosis (>100,000/µL)</td><td>platelets, microcytosis, cryogioouiins leukocytosis (lymphocytes >100,000/µL), lipemia, abnormal protein</td></tr><tr><th>Interfering substances: Differential</th><th>lyse-resistant RBCs</th><th>lyse-resistant RBCs</th><th>lyse-resistant RBCs</th></tr><tr><td>Maximum CBCs per hour/Maximum CBCs and differentials per hour Minimum specimen volume open/Closed/Sample dead volume closed</td><td>80/80 85 μL/150 μL/1 mL</td><td>80/80 85 μL/150 μL/1 mL</td><td>100/100 85 μL/150 μL/1 mL</td></tr><tr><td>Microsample capability Prepares microscope slides automatically or flags problems for slide prep Number of automatic slidemakers available/List price</td><td>yes no </td><td>yes no —</td><td>yes no</td></tr><tr><td>Archives patient data/Previous patient results incl. with recent results Maximum archived data accessible when system online No. specimens for which numeric results saved in memory at once No. specimens for which histo/cytogram results saved in memory at once</td><td>yes/yes 10,000 samples 10,000 samples 10,000 samples</td><td>yes/yes 10,000 samples 10,000 samples 10,000 samples</td><td>yes/yes 10,000 samples 10,000 samples 10,000 samples</td></tr><tr><td>Performs delta checks Tags and holds results for followup, confirmatory testing, or rerun Parameters for flags for holding samples defined by user or vendor Scattergram display: cell-specific color Histogram display: color with thresholds User interface can display choice of specimen/result information</td><td>yes yes user or vendor yes yes yes</td><td>yes yes user or vendor yes yes yes</td><td>yes yes yes yes yes yes</td></tr><tr><td>LIS interface formats supported Information transferred on LIS interface</td><td>RS-232C/TCP-IP, ASTM numeric and flag results, histograms and scatterplots, patient demographics, orders</td><td>RS-232/TCP-IP, ASTM numeric and flag results, histograms and scatterplots, patient demographics, orders</td><td>ASTM numeric and flag results, histograms and scatterplots, instrument to LIS; patient demographics, orders, LIS to instrument—</td></tr><tr><td>LOINC codes transmitted with all results/Sent in message to LIS/ Listing of machine codes and corresponding LOINC for each test Interface available or planned to automated specimen-handling system</td><td>no/no/yes </td><td>no/no/yes </td><td>broadcast; host query for demographics and orders no/no/yes</td></tr><tr><td>Bar-code symbologies read on specimen tube</td><td>Codabar, codes 39 and 128, Interleaved 2 of 5, ITF, NW7, EAN 8 and 13</td><td>Codabar, codes 39 and 128, Interleaved 2 of 5, ITF, NW7, EAN 8 and 13</td><td>Codabar, codes 39 and 128, ASTM, Interleaved 2 of 5, ITF, NW7</td></tr><tr><td>Time required for maintenance by lab personnel</td><td>daily: <3 minutes</td><td>daily: <3 minutes</td><td>daily: <3 minutes</td></tr><tr><td>Onboard diagnostics for troubleshooting/Limited to software problems Manufacturer can perform diagnostics via modem</td><td>yes/no yes, also via Internet</td><td>yes/no yes, also via Internet</td><td>yes/no yes</td></tr><tr><td>Distinguishing features (supplied by company)</td><td>remote diagnostics; online QC; random access; discrete testing; reagent monitoring; chartable report formats; unique specimen-gating, software is FDA Part II compliant; body fluids now FDA cleared; standardized technology, reagents, controls, and operations with other X series analyzers; XT-V for use in toxicology, research, and veterinary reference labs</td><td>high throughput, remote diagnostics; online QC; random access; fluorescent optical platelets; discrete testing; reagent monitoring; customized chartable report formats; body fluids, standardized technology, reagents, controls, and operations with other X series analyzers; IG # & %, RET-He; XT-V unit for use in toxicology, research, and veterinary reference labs</td><td>unique testing parameters: fluorescent optical platelets, IG #&%, RET-He, body fluids (CSF, serous, synovial), WBC/RBC/TC and two-part differential; standardized technology, reagents, controls, and operations with other X series analyzers; simplified operations with extended linearities, high- throughput, remote monitoring capabilities</td></tr></tbody></table>		

Note: a dash in lieu of an answer means company did not answer question

Hematology analyzers

Part 9 of 11 See captodayonline.com/productguides	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic Road Lincolnshire, IL 60069	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic Road Lincolnshire, IL 60069 900 270 7620, www.acamay.com/up	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic Road Lincolnshire, IL 60069
Name of instrument	Sysmex XE-2100D	Sysmex XE-2100	Sysmex XE-5000
First year installed in U.S./Outside U.S./No. of units sold in 2011 No. units installed in U.S./Outside U.S./List price	2004/2004/>15 >200/>205/\$235,640	- 1999/—/~65 1,325/>5,000/\$248,251	- 2008/2008/>80 >1,000/>3,500/\$270,424
Test menu: • Chartable (standard menu: WBC, RBC, Hb, Hct, MCV, MCH, MCHC, PLT, %&# neut, mono, lymph, eos, baso):	standard menu (left) plus: RDW-SD, RDW-CV	standard menu (left) plus: NRBC %&#, retic %&#, RDW-SD, RDW-CV, IRF, PLT-O, HPC#, MPV, IG%, IG#,</td><td>standard menu (left) plus: NRBC %&#, retic %&#, RDW-SD, RDW-CV, IRF, PLT-O, HPC#, MPV, IG%, IG#,</td></tr><tr><td>Laboratory</td><td>_</td><td>RET-He, IPF</td><td>RET-He, IPF</td></tr><tr><td>• Flags</td><td>PLT clumps, PLT abnormal distribution, WBC abnormal scattergram, blast, left shift, atypical lymphocytes, abnormal lymphocytes/blast, RBC abnormal distribution, RBC lyse resistance, RBC agglutinins, turbidity</td><td>PLT clumps, RBC agglut, turbidity, WBC abnormal scattergram, RBC abnormal distribution, PLT abnormal distribution, RBC lyse resistance, blasts, left shift, atypical lymphocytes, abnormal lymphocytes/blast., reticulocyte ABN scattergram</td><td>PLT clumps, PLT ABN distribution, WBC ABN scattergram, blast, left shift, atypical lymphocytes, abnormal lymphocytes/blast, RBC ABN distribution, RBC lyse resistance, RBC agglutinins, turbidity</td></tr><tr><td>FDA-cleared tests not clinically released Tests not available but submitted for 510(k) clearance</td><td>_</td><td>_</td><td>_</td></tr><tr><td>Tests in development Tests for research use only</td><td>— P-LCR. PCT. PDW</td><td>— P-LCR. PCT. PDW</td><td>_</td></tr><tr><td>Tests unique to analyzer</td><td>optional: IG% & IG#</td><td>HPC#, IG%, IG#, RET He, IPF</td><td>reticulocyte hemoglobin, immature platelet fraction, hematopoietic progenitor cell, immature reticulocyte fraction, reportable immature granulocyte #&%, RBC/WBC/TC/two-part differential</td></tr><tr><td>Differential method(s) used</td><td>fluorescent flow cytometry</td><td>fluorescent flow cytometry, RF/DC detecting method</td><td>fluorescent flow cytometry, RF/DC detection method</td></tr><tr><td>Linearity: • WBC count/RBC count • Hemoglobin/Platelet • MCV (fL) or Hct (%) Precision: • WBC count/RBC count • Hemoglobin/Platelet • MCV or Hct Accuracy of automated differential compared with manual differential (per CLSI H-20A2)</td><td>$\begin{array}{l} 0-440/0-8\\ 0-25/0-5,000\\ 0-75 \ (Hct)\\ \leq 3 \ percent/\leq 1.5 \ percent\\ \leq 1.0 \ percent/\leq 4.0 \ percent\\ \leq 1.5 \ percent \ (Hct)\\ neut\% \ r=0.95, \ y=0.92x+5.46; \ lymph\% \ r=0.95,\\ y=0.88x+2.46; \ mon0\% \ r=0.79, \ y=0.77x+1.88; \ eos\%\\ r=0.92, \ y=0.97x+0.29; \ baso\% \ r=0.82, \ y=1.01x+0.01;\\ NRBC\% \ r=0.96, \ y=1.12x+0.11; \ IG\% \ r=0.83,\\ y=0.9332x+0.0922 \end{array}$</td><td>0-440/0-8 0-25/0-5,000 0-75 (Hct) <3 percent/<1.5 percent <1.0 percent/<4.0 percent <1.5 percent (Hct) neut% r=0.95, y=0.92x+5.46; lymph% r=0.95, y=0.88x+2.46; mono% r=0.79, y=0.77x+1.88; eos% r=0.92, y=0.97x+0.29; baso% r=0.82, y=1.01x+0.01; NRBC% r=0.96, y=1.12x+0.11; IG% r=0.83, y=0.9332x+0.0922</td><td>0-440/0-8 0-25/0-5,000 0-75 (Hct) <3 percent/<1.5 percent <1.0 percent/<4.0 percent <1.5 percent (Hct) neut% r=0.95, y=0.92x+5.46; lymph% r=0.95, y=0.88x+2.46; mono% r=0.79, y=0.77x+1.88; eos% r=0.92, y=0.97x+0.29; baso% r=0.82, y=1.01x+0.01; NRBC% r=0.96, y=1.12x+0.11; IG% r=0.83, y=0.9332x+0.0922</td></tr><tr><td>Interfering substances: • WBC</td><td>cold agglutinins, PLT aggregation, cryoglobulin, lyse-resistant RBCs, NRBCs</td><td>cold agglutinin, PLT aggregation, nucleated RBCs, cryoglobulin, lyse-resistant RBCs</td><td>cold agglutinins, PLT aggregation, nucleated RBCs, cryoglobulin, lyse-resistant RBCs</td></tr><tr><td>• RBC</td><td>cold agglutinins, severe microcytosis, fragmented RBCs, leukocytosis</td><td>cold agglutinins, severe microcytosis, fragmented RBCs, large No. giant PLTs, in vitro hemolysis</td><td>cold agglutinins, severe microcytosis, fragmented RBCs, large number giant PLTs, in vitro hemolysis</td></tr><tr><td>• MCV or Hct</td><td>Hct: cold agglutinins, ABN red cell fragility, spherocytosis, leukocytosis</td><td>Hct: cold agglutinins, leukocytosis, ABN red cell fragility, spherocytosis</td><td>Hct: cold agglutinins, leukocytosis, ABN red cell fragility, spherocytosis</td></tr><tr><td>Platelet</td><td>pseudothrombocytopenia, PLT aggregation, increased microcytosis, megaloblasts</td><td>pseudothrombocytopenia, PLT aggregation, increased microcytosis, megalocytic PLTs</td><td>pseudothrombocytopenia, PLT aggregation, increased microcytosis, megalocytic PLTs</td></tr><tr><td>• Hemoglobin</td><td>lipemia, ABN proteins, leukocytosis (>100,000/µL)</td><td>lipemia, ABN proteins, leukocytosis (>100,000/µL)</td><td>lipemia, ABN proteins, leukocytosis (>100,000/µL)</td></tr><tr><td>Interfering substances: Differential</td><td>lyse-resistant RBCs</td><td>lyse-resistant RBCs</td><td>lyse-resistant RBCs</td></tr><tr><td>Maximum CBCs per hour/Maximum CBCs and differentials per hour Minimum specimen volume open/Closed/Sample dead volume closed</td><td>150/150 130 μL/200 μL/1 mL</td><td>150/150 130 μL/200 μL/1 mL</td><td>150/150 130 μL/200 μL/1 mL</td></tr><tr><td>Microsample capability Prepares microscope slides automatically or flags problems for slide prep</td><td>yes yes (with Alpha or HST upgrade)</td><td>yes yes (with Alpha or HST upgrade)</td><td>yes yes (with Alpha or HST upgrade)</td></tr><tr><td>Number of automatic slidemakers available/List price</td><td>>1,000/—</td><td>>1,000/—</td><td>>1,200/—</td></tr><tr><td>Archives patient data/Previous patient results incl. with recent results Maximum archived data accessible when system online No. specimens for which numeric results saved in memory at once No. specimens for which histo/cytogram results saved in memory at once</td><td>yes/yes 10,000 samples 10,000 samples 10,000 samples</td><td>yes/yes 10,000 samples 10,000 samples 10,000 samples</td><td>yes/yes 10,000 samples 10,000 samples 10,000 samples</td></tr><tr><td>Performs delta checks Tags and holds results for followup, confirmatory testing, or rerup</td><td>yes ves</td><td>yes ves</td><td>yes ves</td></tr><tr><td>Parameters for flags for holding samples defined by user or vendor</td><td>user or vendor</td><td>user or vendor</td><td>yes</td></tr><tr><td>Histogram display: color with thresholds</td><td>yes ves</td><td>yes ves</td><td>yes ves</td></tr><tr><td></td><td>DC-2220//TCD ID</td><td>DC-2220/JCD ID</td><td>ACTM 1204 TCD ID ACTM 51201</td></tr><tr><td>Information transferred on LIS interface</td><td>numeric and flag results, histograms and scatterplots, patient demographics, orders</td><td>numeric and flag results, histograms and scatterplots, patient demographics, orders</td><td>numeric and flag results, histograms and scatterplots, instrument to LIS; patient demographics, orders, LIS to instrument—</td></tr><tr><td>LOINC codes transmitted with all results/Sent in message to LIS/</td><td>no/no/yes</td><td>no/no/yes</td><td>no/no/yes</td></tr><tr><td>Listing of machine codes and corresponding LOINC for each test Interface available or planned to automated specimen-handling system</td><td>on automation platform</td><td>on automation platform</td><td>Roche Diagnostics, and Labotix, A & T, Thermo, IDS</td></tr><tr><td>Bar-code symbologies read on specimen tube</td><td>Codabar, codes 39 and 128, ASTM, Interleaved 2 of 5, ITF, NW7, EAN 8 and 13, ISBT</td><td>Codabar, codes 39 and 128, Interleaved 2 of 5, ITF, NW7, EAN 8 and 13</td><td>Codabar, codes 39 and 128, ASTM, Interleaved 2 of 5, ITF, NW7, EAN 8 and 13</td></tr><tr><td>Accommodates bar-code placement per CLSI standard Auto2A</td><td>yes</td><td>yes</td><td>_</td></tr><tr><td>Time required for maintenance by lab personnel Onboard diagnostics for troubleshooting/Limited to software problems Manufacturer can perform diagnostics via modem</td><td>daily: <3 minutes yes/no yes, also via Internet</td><td>daily: <3 minutes yes/no yes, also via Internet</td><td>daily: <3 minutes yes/no yes, also via Internet</td></tr><tr><td>Distinguishing features (supplied by company) Note: a dash in lieu of an answer means company did not answer question or question is not applicable</td><td>150 CBCs per hour; platelet linearity to 5 million, hematocrit extended to 75 percent; standardized technology, reagents, controls and operations; ISBT-compliant; FDA-cleared application for blood component products in specified anticoagulants</td><td>throughput of 150 CBCs per hour; random access; discrete testing; online QC; remote diagnostics, body fluid analysis; platelet linearity to 5 million, hematocrit linear to 75 percent; hematopoietic progenitor cell testing; immature granulocyte enumeration; immature platelet fraction; reticulocyte hemoglobin equivalent; standardized reagents, controls, and operations with other Sysmex X series analyzers</td><td>low-end linearity for all body fluids; two- part differential (mono nuclear % + # and polymorphonuclear % + # or body fluid; reticulocyte hemoglobin content; immature platelet fractions; throughput of 150 CBCs per hour; random access; discrete testing; online QC; remote diagnostics, body fluid analysis; platelet linearity to 5 million, hematocrit linear to 75 percent; hematopoietic progenitor cell testing; immature granulocyte enumeration; immature platelet fraction; reticulocyte hemoglobin equivalent; standardized reagents, controls, and operations with other X series analyzers</td></tr></tbody></table>	

Hematology analyzers

Part 10 of 11	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic Road	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic Road	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic Road
See captodayonline.com/productguides for an interactive version of guide	Lincolnshire, IL 60069 800-379-7639 www.sysmex.com/us	Lincolnshire, IL 60069 800-379-7639 www.sysmex.com/us	Lincolnshire, IL 60069 800-379-7639 www.sysmex.com/us
Name of instrument	Sysmex XE-Alpha N/HST-N	Sysmex XN-1000	Sysmex XN-2000
First year installed in U.S./Outside U.S./No. of units sold in 2011 No. units installed in U.S./Outside U.S./List price	2000/—/>100 >1,000/1,400/\$360,000 - \$1,000,000	2012/2011/— 0/80/\$202,667	2012/2011/— 0/44/\$402,667
Test menu: • Chartable (standard menu: WBC, RBC, Hb, Hct, MCV, MCH, MCHC, PLT, %&# neut, mono, lymph, eos, baso):	standard menu (left) plus: NRBC%&#, retic%&#, RDW-SD, RDW-CV, IRF, PLT-0, HPC#, MPV, IG%, IG#, RET-He, IPF</td><td>standard menu (left) plus: NRBC%, NRBC#, IG%, IG#, MPV, PLT-F, IPF, RDW-CV, RDW-SD, Retic#, Retic%, IRF, RET-He; body fluids: RBC-BF, TC-BF, WBC-BF, MN%, MN#, PMN%, PMN#</td><td>standard menu (left) plus: NRBC%, NRBC#, IG%, IG#, MPV, PLT-F, IPF, RDW-CV, RDW-SD, Retic#, Retic%, IRF, RET-He; body fluids: RBC-BF, TC-BF, WBC-BF, MN%, MN#, PMN%, PMN#</td></tr><tr><td>• Laboratory • Flags</td><td> user-defined, all-inclusive</td><td>— blasts/abnormal lymphocytes, left shift, atypical lymphocytes, RBC agglutination, turbidity/HGB interf, iron deficiency, HGB defect, fragments, PLT clumps</td><td>— blasts/abnormal lymphocytes, left shift, atypical lymphocytes, RBC agglutination, turbidity/HGB interf, iron deficiency, HGB defect, fragments, PLT clumps</td></tr><tr><td>FDA-cleared tests not clinically released Tests not available but submitted for 510(k) clearance</td><td></td><td></td><td>_</td></tr><tr><td>Tests in development Tests for research use only Tests unique to analyzer</td><td>— P-LCR, PCT, PDW NRBC, HPC#, IG%, IG#, RET-He, immature platelet function</td><td>— IG%, IG#, PLT-F, IPF, RET-He; body fluids: two-part differential MN%, MN#, PMN%, PMN#</td><td>— IG%, IG#, PLT-F, IPF, RET-He; body fluids: two-part differential MN%, MN#, PMN%, PMN#</td></tr><tr><td>Differential method(s) used</td><td>fluorescent flow cytometry, RF/DC detecting method</td><td>fluorescent flow cytometry with side fluorescent light, forward-scattered light and side-scattered</td><td>fluorescent flow cytometry with side fluorescent light, forward-scattered and side-scattered light</td></tr><tr><td>Linearity: • WBC count/RBC count • Hemoglobin/Platelet • MCV (fL) or Hct (%) Precision: • WBC count/RBC count • Hemoglobin/Platelet • MCV or Hct Accuracy of automated differential compared with manual differential (per CLSI H-20A2)</td><td>0-440/0-8 0-25/0-5,000 0-75 (Hct) <3 percent/<1.5 percent <1.0 percent/<4.0 percent <1.0 percent (Hct) neut% r=0.95, y=0.92x+5.46; lymph% r=0.95, y=0.88x+2.46; mon0% r=0.79, y=0.77x+1.88; eos% r=0.92, y=0.97x+0.29; baso% r=0.82, y=1.01x+0.01; NRBC% r=0.96, y=1.12x+0.11; IG% r=0.83,</td><td>light 0.00-440.00/0.00-8.60 0.0-26.0/0-5,000 0.0-75.0% (HCT) < 3.0%/< 1.5% < 1.0%/< 4.0% HCT < 1.5% —</td><td>0.00-440.00/0.00-8.60 0.0-26.0/0-5,000 0.0-75.0% (HCT) < 3.0%/< 1.5% < 1.0%/< 4.0% HCT < 1.5%</td></tr><tr><td>Interfering substances: • WBC</td><td>y=0.9332x+0.0922 cold agglutinins, PLT aggregation, nucleated RBCs, cryoglobulins, lyse-resistant RBCs</td><td>no significant interference up to: 39.4 mg/dL for bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for hemolysis, 30.320 0D for intralipid, 2880 0D for chyle</td><td>no significant interference up to: 39.4 mg/dL for bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for hemolysis, 30.320 0D for intralipid, 2880 0D for chyle</td></tr><tr><td>• KRC</td><td>cold agglutinins, severe microcytosis, fragmented RBCs, large No. giant PLTs, in vitro hemolysis</td><td>no significant interterence up to: 39.4 mg/oL tor bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for hemolysis, 55.980 0D for intralipid, 2880 0D for chyle</td><td>no significant interterence up to: 39.4 mg/oL tor bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for hemolysis, 55.980 OD for intralipid, 2880 OD for chyle</td></tr><tr><td>• MCV or Hct</td><td>Hct: cold agglutinins, leukocytosis, ABN red cell fragility, spherocytosis</td><td>no significant interference up to: 39.4 mg/dL for bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for hemolysis. 55 980 0D for intralipid. 2880 0D for chyle</td><td>no significant interference up to: 39.4 mg/dL for bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for hemolysis, 55 980 0D for intralipid, 2880 0D for chyle</td></tr><tr><td>Platelet</td><td>pseudothrombocytopenia, PLT aggregation, increased microcytosis, megalocytic PLTs</td><td>no significant interference up to: 39.4 mg/dL for bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for bemolusie 55.980 0D for intralinid 2880 0D for chule</td><td>no significant interference up to: 39.4 mg/dL for bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for benadusia 55.980 0D for intralinid 2880 0D for chule</td></tr><tr><td>• Hemoglobin</td><td>lipemia, ABN proteins, leukocytosis (>100,000/µL)</td><td>no significant interference up to: 39.4 mg/dL for bilirubin C, 37.4 mg/dL for bilirubin F, 199 mg/dL for hemolysis</td><td>no significant interference up to: 39.4 mg/dL for bilirubin C, 37.4 mg/dL for bilirubin F, 199 mg/dL for hemolysis</td></tr><tr><td>Interfering substances: Differential</td><td>lyse-resistant RBCs</td><td></td><td></td></tr><tr><td>Maximum CBCs per hour/Maximum CBCs and differentials per hour Minimum specimen volume open/Closed/Sample dead volume closed Microsample capability Prepares microscope slides automatically or flags problems for slide prep Number of automatic slidemakers available/List price</td><td>150/150 per analyzer on automation system 130 μL/200 μL/1 mL yes yes >1,700/\$250,000</td><td>100/100 88 μL/88 μL/1 mL yes yes —</td><td>200/200 88 μL/88 μL/1 mL yes yes —</td></tr><tr><td>Archives patient data/Previous patient results incl. with recent results Maximum archived data accessible when system online No. specimens for which numeric results saved in memory at once No. specimens for which histo/cytogram results saved in memory at once</td><td>yes/yes 10,000 samples 10,000 samples; 20,000 orders 10,000 samples; two years plus, with optional decision logic software</td><td>yes/yes 100,000 100,000 100,000</td><td>yes/yes 100,000 100,000 100,000</td></tr><tr><td>Performs delta checks Tags and holds results for followup, confirmatory testing, or rerun Parameters for flags for holding samples defined by user or vendor Scattergram display: cell-specific color Histogram display: color with thresholds User interface can display choice of specimen/result information</td><td>yes yes user and vendor yes yes yes</td><td>yes yes yes yes yes yes</td><td>yes yes yes yes yes yes</td></tr><tr><td>LIS interface formats supported</td><td>RS-232C/TCP IP</td><td>[XN series ASTM1381-95/ASTM1894-97] or [XN</td><td>[XN series ASTM1381-95/ASTM1894-97] or [XN</td></tr><tr><td>Information transferred on LIS interface</td><td>numeric and flag results, histograms and scatterplots, patient demographics, orders</td><td>series ASTM1381-02/ASTM1394-97J numeric and flag results, histograms and scatterplots, instrument to LIS; patient demographics, orders, LIS to instrument— broadcast bact upper for demographics and orders</td><td>series ASTM1381-U2/ASTM1394-97J numeric and flag results, histograms and scatterplots, instrument to LIS; patient demographics, orders, LIS to instrument— brandpact bact upper for demographics and orders</td></tr><tr><td>LOINC codes transmitted with all results/Sent in message to LIS/ Listing of machine codes and corresponding LOINC for each test Interface available or planned to automated specimen-handling system Bar-code symbologies read on specimen tube</td><td>no/no/yes Roche, Labotix, IDS, A&T, Thermo engen Codabar. codes 39 and 128, Interleaved 2 of 5, ITF,</td><td>no/no/yes none Codabar. codes 39 and 128, ITF, NW7, ISBT 128, JAN/</td><td>none Codabar, codes 39 and 128, ITF, NW7, ISBT 128, JAN/</td></tr><tr><td>Accommodates bar-code placement per CLSI standard Auto2A</td><td>NW7, EAN 8 and 13 yes</td><td>EAN/UPC yes</td><td>EAN/UPC yes</td></tr><tr><td>Time required for maintenance by lab personnel Onboard diagnostics for troubleshooting/Limited to software problems Manufacturer can perform diagnostics via modem</td><td>daily: <3 minutes (operator time) yes/no yes, also via Internet</td><td>daily: <1 minute (operator time) yes/no yes</td><td>daily: <1 minute (operator time) yes/no yes</td></tr><tr><td>Distinguishing features (supplied by company)</td><td>high-throughput, flexible, scalable configurations (>125 standard configurations available); platelet linearity—5 million; new parameters for platelet monitoring—IPF and reticulocyte Hb measurement and RET-He, hematopoietic progenitor cell analysis, lavender top management, standardized technology, reagents, controls, and operations; broad clinical reportable ranges; enhanced clinical parameters to support preventive care and disease management</td><td>reportable parameters include IG %/#, RET- He, fluorescent PLT, body fluid with two-part differential; onboard pre-loaded decision rules including automated rerun-reflex capabilities; optional wagons for complete reagent management and option of using a concentrated reagent</td><td>fully integrated co-primary hematology solution consisting of two analytical modules connected with a single sampler, providing maximum productivity and efficiency with workload balancing; reportable parameters, include IG %/#, RET-He, fluorescent PLT, body fluid with two-part differential, onboard pre-loaded decision rules including automated rerun-reflex capabilities; optional wagons for complete reagent management and option of using a concentrated reagent</td></tr></tbody></table>		

Note: a dash in lieu of an answer means company did not answer question

Hematology analyzers

		1
Part 11 of 11 See captodayonline.com/productguides for an interactive version of guide	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic, Lincolnshire, IL 60069 800-379-7639 www.sysmex.com/usa	Sysmex America Tammy Kutz communications@sysmex.com 577 Aptakisic, LincoInshire, IL 60069 800-379-7639 www.sysmex.com/usa
Name of instrument First year installed in U.S./Outside U.S./No. of units sold in 2011 No. units installed in U.S./Outside U.S./List price	Sysmex XN-3000 2012/2011/— 0/32/\$562,667	Sysmex XN-9000 2012/2011/— 0/38/varies based on configuration
Test menu: • Chartable (standard menu: WBC, RBC, Hb, Hct, MCV, MCH, MCHC, PLT, %&# neut, mono, lymph, eos, baso):	standard menu (left) plus: NRBC%, NRBC#, IG%, IG#, MPV, PLT-F, IPF, RDW-CV, RDW-SD, Retic#, Retic%, IRF, RET-He; body fluids: RBC- BF, TC-BF, WBC-BF, MN%, MN#, PMN%, PMN#	standard menu (left) plus: NRBC%, NRBC#, IG%, IG#, MPV, PLT-F, IPF, RDW-CV, RDW-SD, Retic#, Retic%, IRF, RET-He; body fluids: RBC- BF, TC-BF, WBC-BF, MN%, MN#, PMN%, PMN#
• Laboratory • Flags	blasts/abnormal lymphocytes, left shift, atypical lymphocytes, RBC agglutination, turbidity/HGB interf, iron deficiency, HGB	
FDA-cleared tests not clinically released Tests not available but submitted for 510(k) clearance	— — —	— — —
Tests for research use only Tests unique to analyzer		 IG%, IG#, PLT-F, IPF, RET-He; body fluids: two-
Differential method(s) used	part differential MN%, MN#, PMN%, PMN#	part differential MN%, MN#, PMN%, PMN#
Linearity: • WBC count/RBC count	fluorescent light, forward-scattered and side- scattered light 0.00-440.00/0.00-8.60	fluorescent light, forward-scattered and side- scattered light 0.00-440.00/0.00-8.60
Hemoglobin/Platelet MCV (fL) or Hct (%) Precision: WBC count(RBC count	0.0-26.0/0-5,000 0.0-75.0% (HCT) < 3.0%/< 1.5%	0.0-26.0/0-5,000 0.0-75.0% (HCT) < 3.0%/c 1.5%
Hemoglobin/Platelet MCV or Hct	< 1.0%/< 4.0% HCT < 1.5%	< 1.0%/< 4.0% HCT < 1.5%
Accuracy of automated differential compared with manual differential (per CLSI H-20A2) Interfering substances: • WBC		
	for bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for hemolysis, 30.320 OD for intralipid,	for bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for hemolysis, 30.320 OD for intralipid,
• RBC	2880 OD for chyle no significant interference up to: 39.4 mg/dL for bilirubin C. 37.4 mg/dL for bilirubin F. 996	2880 OD for chyle no significant interference up to: 39.4 mg/dL for bilirubin C. 37.4 mg/dL for bilirubin F. 996
	mg/dL for hemolysis, 55.980 OD for intralipid, 2880 OD for chyle	mg/dL for hemolysis, 55.980 OD for intralipid, 2880 OD for chyle
MCV or Hct	no significant interference up to: 39.4 mg/dL for bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for hemolysis, 55.980 OD for intralipid.	no significant interference up to: 39.4 mg/dL for bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for hemolysis. 55.980 OD for intralipid.
• Platelet	2880 0D for chyle no significant interference up to: 39.4 mg/dL	2880 0D for chyle no significant interference up to: 39.4 mg/dL
	for bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for hemolysis, 55.980 OD for intralipid, 2880 OD for chvle	for bilirubin C, 37.4 mg/dL for bilirubin F, 996 mg/dL for hemolysis, 55.980 OD for intralipid, 2880 OD for chvle
• Hemoglobin	no significant interference up to: 39.4 mg/dL for bilirubin C, 37.4 mg/dL for bilirubin F, 199 mg/dl_for_homolycic	no significant interference up to: 39.4 mg/dL for bilirubin C, 37.4 mg/dL for bilirubin F, 199 mg/dL for homolycic
Interfering substances: Differential	— —	mg/aL for nemolysis —
Maximum CBCs per hour/Maximum CBCs and differentials per hour Minimum specimen volume open/Closed/Sample dead volume closed Microscample canability	200/200 88 µL/88 µL/1 mL	200/200 88 µL/88 µL/1 mL ves
Prepares microscope slides automatically or flags problems for slide prep Number of automatic slidemakers available/List price	yes —/included	yes —/included
Archives patient data/Previous patient results incl. with recent results Maximum archived data accessible when system online	yes/yes 100,000 100,000	yes/yes 100,000 100,000
No. specimens for which histo/cytogram results saved in memory at once Performs delta checks	100,000 yes	100,000 yes
Tags and holds results for followup, confirmatory testing, or rerun Parameters for flags for holding samples defined by user or vendor	yes yes	yes yes
Scattergram display: cell-specific color Histogram display: color with thresholds User interface can display choice of specimen/result information	yes yes ves	yes yes ves
LIS interface formats supported	[XN series ASTM1381-95/ASTM1894-97] or	[XN series ASTM1381-95/ASTM1894-97] or
Information transferred on LIS interface	[XN series ASTM1381-02/ASTM1894-97] numeric and flag results, histograms and scatterplots, instrument to LIS; patient demo- graphics, orders, LIS to instrument—broad- cast; host query for demographics and orders	[XN series ASTM1381-02/ASTM1894-97] numeric and flag results, histograms and scatterplots, instrument to LIS; patient demo- graphics, orders, LIS to instrument—broad- cast; host query for demographics and orders
LOINC codes transmitted with all results/Sent in message to LIS/ Listing of machine codes and corresponding LOINC for each test Interface available or planned to automated specimen-handling system Bar-code symbologies read on specimen tube	no/no/yes none Codabar codes 39 and 128 ITE NW7 ISBT	no/no/yes none Codabar codes 39 and 128 ITE NW7 ISBT
Accommodates bar-code placement per CLSI standard Auto2A	128, JAN/EAN/UPC yes	128, JAN/EAN/UPC yes
Time required for maintenance by lab personnel Onboard diagnostics for troubleshooting/Limited to software problems Manufacturer can perform diagnostics via modem	daily: <3 minutes (operator time) yes/no yes	daily: <3 minutes (operator time) yes/no yes
Distinguishing features (supplied by company)	fully integrated co-primary hematology solution consisting of two analytical	fully integrated, scalable automation, configures up to nine modules including
Note: a dash in lieu of an answer means comnany did not answer question	modules connected with a single sampler, plus a fully integrated slidemaker-stainer (SP-10) providing maximum productivity and efficiency with workload balancing; reportable parameters include IG %/#, RET-He, fluorescent PLT, body fluid with two-part differential, onboard pre-loaded decision rules including automated rerun-reflex capabilities; optional wagons for complete reagent management and option of using a concentrated reagent	up to two SP-10 slidemaker/stainers; when coupled with tube sorting and archiving, HbA1c automation and powered by Intel- ligent Automation from the Sysmex WAM, carries the Lavender Top Management capabilities, which manages more than 90 percent of lavender top tubes; concentrated reagents are standard, saving space and reagent changes with a 25× concentrated Cellpack DST diluent; reportable parameters include IG %/#, RET-He, fluorescent PLT, body fluid with two-part differential; on- hoard nre-loaded decision rules including
or question is not applicable		automated rerun/reflex capabilities