

# Role of Dimerization of the Membrane-associated Growth Factor Kit Ligand in Juxtacrine Signaling: The $Sl^{7H}$ Mutation Affects Dimerization and Stability—Phenotypes in Hematopoiesis

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## Summary

The Kit ligand (KL)/Kit receptor pair functions in hematopoiesis, gametogenesis, and melanogenesis. KL is encoded at the murine *steel* (*Sl*) locus and encodes a membrane growth factor which may be proteolytically processed to produce soluble KL. The membrane-associated form of KL is critical in mediating Kit function in vivo. Evidence for a role of cytoplasmic domain sequences of KL comes from the  $Sl^{7H}$  mutation, a splice site mutation that replaces the cytoplasmic domain with extraneous amino acids. Using deletion mutants and the  $Sl^{7H}$  allele, we have investigated the role of the cytoplasmic domain sequences of KL in biosynthetic processing and cell surface presentation. The normal KL protein products are processed for cell surface expression, where they form dimers. Both  $Sl^{7H}$  and the cytoplasmic deletion mutants of KL were processed to the cell surface; however, the rate of transport and protein stability were affected by the mutations. Deletion of cytoplasmic domain sequences of KL did not affect dimerization of KL. In contrast, dimerization of the  $Sl^{7H}$  protein was reduced substantially. In addition, we have characterized the hematopoietic cell compartment in  $Sl^{7H}$  mutant mice. The  $Sl^{7H}$  mutation has only minor effects on hematopoiesis. Tissue and peritoneal mast cell numbers were reduced in mutant mice as well as in myeloid progenitors. Interestingly, long-term bone marrow cultures from  $Sl^{7H}$  mice did not sustain the long-term production of hematopoietic cells. In addition, homing of normal hematopoietic progenitors to the spleen of irradiated  $Sl^{7H}/Sl^{7H}$  recipient mice was diminished in transplantation experiments, providing evidence for a role of Kit in homing or lodging. These results demonstrate that the membrane forms of KL exist as homodimers on the cell surface and that dimerization may play an important role in KL/Kit-mediated juxtacrine signaling.

Polypeptides which transmit extracellular signals are most often secreted as soluble factors. An increasing number of growth factors that derive from membrane-anchored precursors and growth factors which function both as soluble and as membrane growth factors have been described in the past several years. They include members of the epidermal growth factor family, e.g., epidermal growth factor, TGF- $\alpha$ , and c-erb/HER2 ligand (1), several hematopoietic growth factors, including the c-kit ligand (KL),<sup>1</sup> CSF-1, IL-1, and TNF (2, 3). Transmembrane

growth factors have also been identified in the nematode *Caenorhabditis elegans* (lin-3) (4) and in *Drosophila*, *bride of sevenless* (*boss*) (5, 6). Several of these transmembrane growth factors release soluble growth factors by proteolytic processing or shedding (7–9). The nonprocessed membrane-anchored growth factors themselves possess biological activities and promote cell adhesion and juxtacrine stimulation in adjacent cells expressing cognate receptors on their cell surface (3). Furthermore, the recently described ligands of the eph-family receptor tyrosine kinases with roles in axon guidance are obligate membrane growth factors (10).

The protooncogene *c-kit* encodes a receptor tyrosine kinase which is a member of the platelet-derived growth factor receptor subfamily (11). *c-kit* and its cognate ligand, KL, also called steel factor, are encoded at the murine *white-*

<sup>1</sup>Abbreviations used in this paper: BFU-E, burst-forming unit-erythroid; BMMC, bone marrow-derived mast cell; CFU-GEMM, CFU-granulocyte erythroid megakaryocyte macrophage; CFU-S, CFU-spleen; endo, endoglycosidase; ER, endoplasmic reticulum; KL, Kit ligand; *Sl*, *steel*; VLA, very late antigen; *W*, white-spotting; WBC, white blood cell.

spotting (*W*) (12, 13) and *steel* (*Sl*) loci (14–16), respectively. The phenotypes of mice with mutations at the *W* and *Sl* loci imply functions of Kit/KL in germ cell development, melanogenesis, and hematopoiesis (17–21). In target cells, KL promotes cell proliferation, survival, cell adhesion/migration, and differentiation. The *steel* gene encodes two KL protein products, KL-1 and KL-2, produced by alternative splicing (2, 22). Both the KL-1 and KL-2 proteins are synthesized as transmembrane proteins and are expressed on the cell surface (2). Both membrane-bound KL-1 and KL-2 can mediate direct cell–cell contact with *c-kit*-expressing cells, e.g., bone marrow–derived mast cells (BMMCs) (22, 23) and primordial germ cells (24, 25). Proteolytic processing of both membrane-bound KL-1 and KL-2 releases biologically active soluble KL proteins, although with different efficiencies and through distinct proteolytic cleavage mechanisms (2, 9, 26). Molecular characterization of various *Sl* alleles has provided important insight into the role of the soluble and membrane-anchored forms of KL (14, 15, 22, 27). Homozygotes for the *Steel-Dickie* mutation (*Sl<sup>d</sup>*) are viable and less severely affected, implying some residual functional activity of KL, but they display all of the pleiotropic effects normally associated with *steel* mutations (17, 18). Molecular analysis showed that the *Sl<sup>d</sup>* allele arose as a result of an intragenic deletion including the transmembrane domain and COOH terminus, generating a secreted KL protein product with normal biological activity (2, 22, 27). The biological characteristics of mice carrying the *Sl<sup>d</sup>* mutation imply that the *Sl<sup>d</sup>* KL protein sustains some activity but is largely defective in facilitating proliferation and survival of target cells, indicating that the membrane-anchored forms of KL play pivotal roles in *c-kit* function. The *Sl<sup>7H</sup>* allele contains a splice donor site mutation resulting in the substitution of amino acids 239–273 in the cytoplasmic domain with 27 extraneous amino acids (28). The phenotypes of mice carrying the *Sl<sup>7H</sup>* allele include white spotting with residual pigmentation on the ears and macrocytic anemia (28, 29). Interestingly, the *Sl<sup>7H</sup>* allele affects male but not female fertility.

The importance of the cell membrane-anchored forms of KL revealed by the phenotypes of the *Sl<sup>d</sup>* mutation and the intriguing phenotypes of mice carrying the *Sl<sup>7H</sup>* allele raise questions about the role of the cytoplasmic domain sequences of KL. Comparison of the cytoplasmic domain sequences of murine and human KL reveals 91% shared identity. In contrast, the extracellular domain sequences of murine and human KL are 79% identical. The strong conservation of the cytoplasmic domain sequences of KL throughout evolution suggests a function for these sequences. The cytoplasmic domain sequences of KL may interact with itself or other molecule(s), and these interactions may play a role in processing to the cell surface, cell surface presentation, and juxtacrine signaling. We have investigated the role of cytoplasmic domain sequences of wild-type KL and the *Sl<sup>7H</sup>* protein maturation and cell surface presentation. We have also investigated the characteristics of hematopoietic cells in mice carrying the *Sl<sup>7H</sup>* al-

lele. The results obtained in this study are discussed in the context of mechanisms governing juxtacrine signaling.

## Materials and Methods

**Construction of KL Deletion Mutants.** Deletion mutations in the cytoplasmic domain of KL were created by the primer-directed PCR mutagenesis method. One 5' primer covering the initiation codon of KL (5'-GACGGAAAGGAATACTTCTCTGTGTT-3') and several 3' primers containing a termination codon located at the desired position were used in the PCR reaction. A plasmid containing the KL cDNA clone 19.1.1 (2) was used as template. The 3' primers used to create the following mutants were 5'-CTCTCTAGATTACAACATACTTATCTC-3', for KL-L263; 5'-ATTTCTAGATTAATTAATCTGTATATT-3', for KL-N254; 5'-TATTCTAGAACTGCCCTTGTAAGACT-3', for KL-V248; and 5'-CCTTTCTAGACTTTACTGTTTCTTCTT-3' for KL-Q241. For generation of the internal deletion mutation, KL- $\Delta$ 241/254, and KL-*Sl<sup>7H</sup>*, an M13 phage DNA containing the KL cDNA clone 19.1.1 and the following oligonucleotides were used: for KL- $\Delta$ 241/254, 5'-ATTATCCTCTTCATTCTGTTTCTTCTTCCA-3'; and for KL-*Sl<sup>7H</sup>*, 5'-TCTTTCTGTTGCAACATACTTCCAGTATAAGGCTCC-3'. All mutations were confirmed by sequence analysis. The mutant KL cDNAs were subsequently cloned into pCDM8 for expression in COS-1 cells.

**Transient Expression of KL cDNAs in COS-1 Cells.** COS-1 cells were transfected with the DEAE-dextran method described previously (2, 30) with minor modifications. Briefly, COS-1 cells were grown to subconfluence 1 d before use, trypsinized, and reseeded on 150-mm petri dishes at a density of  $6 \times 10^6$  cells per dish. After 24 h, the cells had reached  $\sim 70\%$  confluence and were transfected with 5  $\mu$ g of plasmid DNA in the presence of 10% DEAE-dextran (Sigma Chemical Co., St. Louis, MO) for 6–12 h. The medium containing the plasmid DNA was removed, and the cells were shocked with 10% DMSO/PBS<sup>++</sup> for 1 min. Residual DMSO was removed by washing the cells with PBS<sup>++</sup> twice. Transfected COS-1 cells were grown in DME containing 10% FCS, 100 mg/ml L-glutamine, and antibiotics.

**Pulse Chase and Immunoprecipitation Analyses of KL Proteins.** At 72 h after transfection, COS-1 cells expressing KL proteins were used for pulse chase experiments. Cells were incubated with methionine-free DME containing 10% dialyzed FCS for 30 min and labeled with <sup>35</sup>S-methionine (DuPont-NEN, Boston, MA) at 0.5 mCi/ml. At the end of the labeling period, the labeling medium was replaced with regular medium containing an excess amount of methionine. Cell lysates and supernatants were collected at the indicated times. Cell lysates were prepared as described previously (31) in 1% Triton X-100, 20 mM Tris (pH 7.5), 150 mM NaCl, 20 mM EDTA, 10% glycerol, and protease inhibitors phenylmethyl sulfonyl chloride (1 mM) and leupeptin (20  $\mu$ g/ml). For immunoprecipitation analysis of KL protein products, equal volumes of cell lysates or supernatants were immunoprecipitated with excess anti-KL antiserum. KL antiserum was obtained by immunization of rabbits with soluble native and recombinant murine KL, and both antisera were used with indistinguishable results. The anti-KL serum was conjugated to protein A-Sepharose (Pharmacia Biotech, Inc., Piscataway, NJ) and washed three times with wash A (0.1% Triton X-100, 20 mM Tris [pH 7.5], 150 mM NaCl, 10% glycerol). Anti-KL serum—protein A-Sepharose conjugate was incubated with supernatants and cell lysates at 4°C for at least 2 h. The immunoprecipitates

were then washed once in wash B (50 mM Tris, 500 mM NaCl, 5 mM EDTA, 0.2% Triton X-100), three times in wash C (50 mM Tris, 150 mM NaCl, 0.1% Triton X-100, 0.1% SDS, 5 mM EDTA), and once in wash D (10 mM Tris, 0.1% Triton X-100). For gel analysis, immunoprecipitates were solubilized in SDS sample buffer by boiling for 5 min, and analyzed by SDS-PAGE (12%) and autoradiography. For endoglycosidase (endo) H treatment, digestion with endo H (Boehringer Mannheim Biochemicals, Indianapolis, IN) was performed by incubating immunoprecipitates for 16 h at 37°C with 5 mU of endo H in 50  $\mu$ l of 0.1 M sodium phosphate buffer, pH 6.1, containing 0.1% Triton X-100, 0.03% SDS, and 20 mM EDTA.

**Immunofluorescence Microscopy.** At 40 h after transfection, COS-1 cells were trypsinized and reseeded on 12-mm round coverslips. After 32 h, the cells were fixed for 15 min at 25°C with 2% (vol/vol) formaldehyde in PBS and rinsed twice with PBS. Cells were then either used directly for immunofluorescence staining or permeabilized by incubating for 15 min at 25°C with 0.1% (wt/vol) saponin in PBS. For staining, cells were incubated at 25°C for 30 min consecutively with 100  $\mu$ l of normal goat serum, anti-KL serum (1:100 dilution) (32), or an ER marker antibody recognizing the signal sequence receptor (1:100 dilution) (33) and FITC-conjugated goat anti-rabbit antibody, sequentially. The unbound antibody was removed by washing in PBS, and the coverslips were mounted on microscope slides using Permount.

**Covalent Cross-linking of KL Protein, Immunoprecipitation, and Western Blot Analysis.** At 72 h after transfection of COS-1 cells expressing KL proteins, the transfected cells were washed with HBSS<sup>++</sup> and then incubated with 1 mmol/liter bis(sulfosuccinimidyl) substrate (BS<sup>3</sup>) (Pierce, Rockford, IL) for 1 h at 37°C. The reaction was terminated with 10 mmol/liter Tris-HCl (pH 7.5) for 10 min. Cell lysates were prepared by incubating equivalent amounts of cell lysate with anti-mouse stem cell factor mAb (2  $\mu$ g per 10<sup>7</sup> cells) (Genzyme Corp., Cambridge, MA) for 3 h at 4°C. Protein G-Sepharose beads (Pharmacia Biotech, Inc.) were used to collect the antigen-antibody complexes. For gel analysis, immunoprecipitates were solubilized in SDS sample buffer by boiling for 5 min, and analyzed by SDS-PAGE (10%). Proteins were electrophoretically transferred onto nitrocellulose membrane (Bio-Rad Laboratories, Hercules, CA). After transfer, the membrane was incubated with 4% skim milk/TBST (50 mmol/liter Tris-HCl, pH 7.5, 150 mmol/liter NaCl, 0.05% Tween 20) for 12 h at room temperature, and Western blot analysis using anti-mouse stem cell factor polyclonal antibody (1:500 dilution; Genzyme Corp.) and enhanced chemiluminescence detection (Pierce Chemical Co., Rockford, IL) were used for identification.

**Mast Cell Adhesion Assay.** Transfected COS cells were grown in 24-well plates (7.5  $\times$  10<sup>4</sup> per well). BMMCs, obtained as described previously (34), were washed three times with RPMI complete medium and incubated for 1 h with transfected COS cells (1.5  $\times$  10<sup>5</sup> cells per well). Nonadherent BMMCs were then removed by washing three times with medium, and the BMMCs attached to the COS cells were counted by visual inspection in the microscope. BMMCs were counted as total number in each well or the number attached to individual COS cells. To compete the adhesion of BMMCs to COS cells, the *c-kit* antagonistic antibody ACK2 (a gift from Dr. Nishikawa, Kyoto University, Kyoto, Japan) was used.

**Analysis of Peripheral Blood Parameters and Hematopoietic Progenitor Assays.** Blood samples for platelet and white blood cell (WBC) count were drawn from the retroorbital plexus or the tail vein with a capillary pipette (Unopette; Becton Dickinson Lab-

ware, Rutherford, NJ). Platelet and WBC numbers were determined using a hemacytometer and phase-contrast microscopy. The hematocrit was determined using heparinized microhematocrit tubes (Fisher Scientific Co., Pittsburgh, PA).

For *in vitro* progenitor assays, bone marrow was harvested, and the cellularity was determined as described previously (35). 10<sup>5</sup> bone marrow cells were resuspended in 1 ml of IMDM/0.8% methylcellulose (Fisher Scientific Co.)/30% FBS (Hyclone Laboratories, Inc., Logan, UT)/0.2 mM hemin (Sigma Chemical Co.) containing 50 ng/ml mouse IL-3 (BioSource International, Camarillo, CA), 3 U/ml human erythropoietin (Amgen, Thousand Oaks, CA), and 20 ng/ml recombinant mouse KL. Cultures were incubated at 37°C in a 5% CO<sub>2</sub> atmosphere. After 7 d of incubation, colonies were scored on the basis of gross morphology as erythroid burst (BFU-E), granulocyte/macrophage (CFU-GM), and mixed colonies (CFU-GEMM).

Splenic CFU (CFU-S)-day 12 were assessed by injection of 10<sup>5</sup> donor bone marrow cells suspended in MEM supplemented with Mops and 10% fetal bovine serum into the lateral tail vein of lethally irradiated (9.5 Gy) recipient mice. Mice were killed 12 d later, their spleens were fixed in Bouin's solution and 10% formalin, and macroscopically visible colonies on the surface of the spleens were enumerated.

For long-term bone marrow cultures, bone marrow cells from one femur were flushed into a T-25 flask in Fisher's medium containing 20% horse serum and dexamethasone. Cultures were incubated at 33°C and semidepopulated at weekly intervals.

**Determination of Mast Cell Numbers in the Skin and Peritoneal Cavity.** For the determination of mast cell numbers in the skin, pieces of dorsal skin were removed from different parts of the back, smoothed onto a piece of thick filter paper to keep them flat, and fixed in Carnoy's solution. Tissues were embedded in paraffin, and 5- $\mu$ m sections were stained with 0.1% acidified toluidine blue (pH 4.0). In skin sections, mast cells between the epithelium and the paniculus carnosus were counted under the microscope. The number of mast cells per centimeter length of skin was determined by dividing the mast cell number by the length of each section of skin counted. For each sample, measurements were made in two separate histological sections and averaged to provide the sample mean.

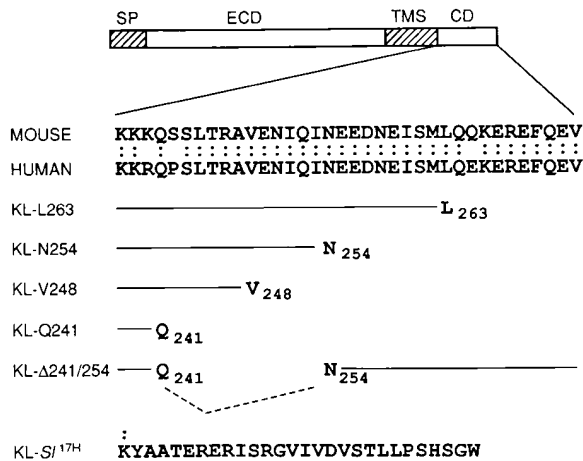
Peritoneal mast cells were obtained from *S<sup>KL2</sup>/S<sup>KL2</sup>* mice and control mice by lavage of the peritoneal cavity with 5 ml PBS. Mast cells were identified by staining with 0.1% toluidine blue (34).

## Results

### *Construction of KL-1 Cytoplasmic Domain Variants and Analysis of Their Turnover Characteristics in COS-1 Cells.*

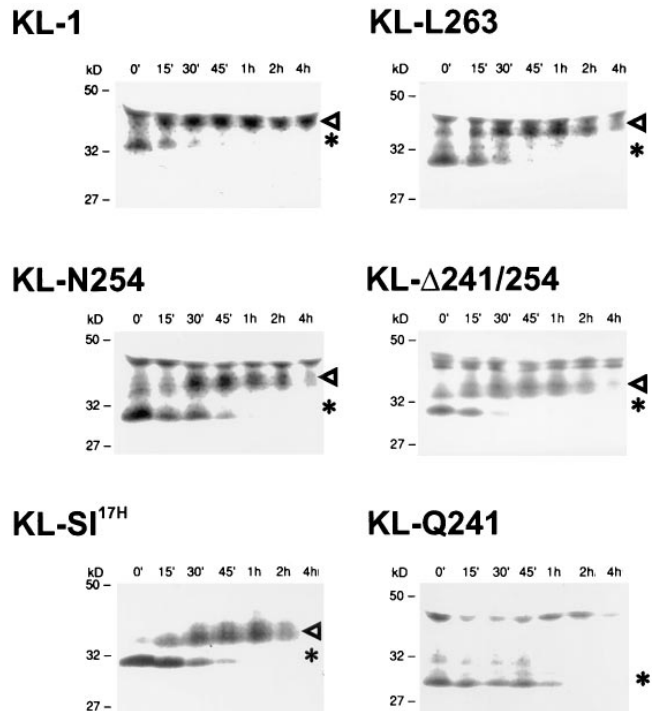
The cytoplasmic domain sequences of KL are highly conserved in evolution. To study roles for cytoplasmic domain sequences of KL, COOH-terminal deletion mutations of varying lengths were constructed (Fig. 1). The wild-type KL-1 cDNA plasmid was used as a template for mutagenesis. The deletion mutations were named according to the position of the COOH-terminal amino acid, e.g., KL-L263, KL-N254, KL-V248, and KL-Q241. In addition, an internal deletion mutation, KL- $\Delta$ 241/254, and KL-S<sup>17H</sup> were constructed. All constructs were subcloned into the eukaryotic expression vector pCDM8 suitable for transient expression in COS-1 cells.

The primary KL translation products are progressively



**Figure 1.** Schematic representation of KL cytoplasmic variants and amino acid sequence of KL-SI<sup>17H</sup>. SP, Signal peptide. ECD, Extracellular domain. TMS, Transmembrane segment. CD, Cytoplasmic domain.

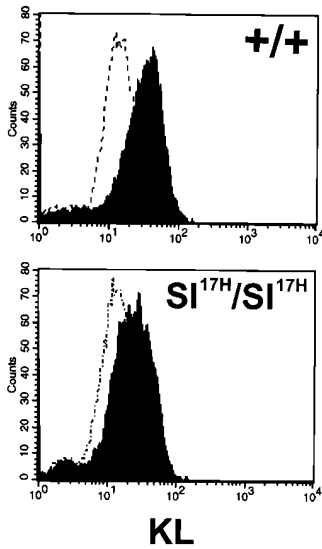
processed primarily by carbohydrate modification as they are transported through the endoplasmic reticulum (ER) and Golgi compartments to the cell surface (2). Biosynthetic processing of the protein products of the KL cytoplasmic domain variants and KL-SI<sup>17H</sup> was investigated by performing pulse chase experiments in transiently transfected COS-1 cells by using <sup>35</sup>S-methionine. Immunoprecipitation analysis using anti-KL antiserum and SDS-PAGE indicated that the biosynthetic processing pattern of all cytoplasmic domain mutants and KL-SI<sup>17H</sup> was similar to that of wild-type KL-1 (Fig. 2). Smaller molecular mass ( $M_r$ ) proteins representing the unglycosylated forms of KL were synthesized: KL-1 (35,000), KL-L263 (33,000), KL-N254 (31,000), KL-V248 (33,000, not shown), KL-Δ241/254 and KL-SI<sup>17H</sup> (33,000), and KL-Q241 (30,000). These proteins were progressively modified by glycosylation to form more mature products representing the cell surface forms. The respective  $M_r$  for the mature proteins were as follows: KL-1, 45,000 daltons; KL-L263, 43,000 daltons; KL-N254, 41,000 daltons; KL-V248 (42,000 daltons, not shown), KL-Δ241/254 and KL-SI<sup>17H</sup> (43,000 daltons); and KL-Q241 (40,000 daltons) (Fig. 2). Previous cell surface I<sup>125</sup>-iodination experiments have shown that the 45-kD KL-1 protein represented the mature KL-1 protein on the cell membrane (2). Quantitation of the different KL protein products during the chase indicated that after 15 min, ~50% of the immature 35-kD KL-1 protein was processed to the mature 45-kD form. With KL-SI<sup>17H</sup>, this process was somewhat delayed, in that 30–45 min was required for 50% of the smaller 33,000- $M_r$  protein products to be processed to their higher 43,000- $M_r$  forms. KL-L263, KL-N254, KL-V248, and KL-Δ241/254 were also delayed in their maturation to a similar extent as KL-SI<sup>17H</sup>. KL-Q241 displayed the most striking delay of maturation, as only a small fraction of the KL-Q241 protein products was processed to the mature 40-kD protein. In agreement with these findings, stromal cells derived from homozygous SI<sup>17H</sup>/SI<sup>17H</sup> mice express a reproducibly reduced level of cell surface KL



**Figure 2.** Biosynthetic characteristics of KL-1 cytoplasmic domain variants and KL-SI<sup>17H</sup>. COS-1 cells were transfected with various mutant constructs, KL-1, KL-L263, KL-N254, KL-Δ241/254, KL-SI<sup>17H</sup>, and KL-Q241, labeled with <sup>35</sup>S-methionine for 30 min, and chased in regular medium. Cell lysates were collected at designated time points, immunoprecipitated with anti-KL antibody, and analyzed by SDS-PAGE (12%). Arrowheads, Mature glycosylated KL proteins. \*Immature forms. Molecular mass ( $M_r$ ) markers are indicated in kilodaltons.

(~60% of normal levels) as determined by FACS<sup>®</sup> analysis compared with the low levels of cell surface KL seen in normal controls (Fig. 3).

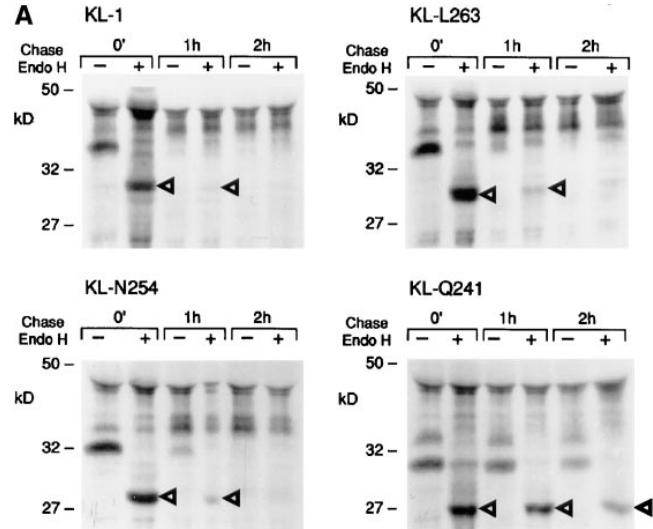
We then determined whether the wild-type KL-1 and the mutant KL protein products follow the same maturation process through the ER and the Golgi complex. Upon reaching the medial Golgi complex, *N*-acetylglucosamine is added to glycoproteins, rendering the carbohydrates resistant to the cleavage by endo H (36, 37). To characterize the maturation pathway of mutant KL proteins, we used sensitivity to endo H treatment. COS-1 cells transfected with various constructs were pulse labeled with <sup>35</sup>S-methionine and then chased in the presence of excess nonradioactive methionine for 0, 1, and 2 h, and KL protein products were immunoprecipitated from cell lysates and digested with endo H. The wild-type KL-1 protein became resistant to endo H treatment at 1 and 2 h of chase, consistent with rapid transit through the ER Golgi complex. The KL-L263 and KL-N254 protein products displayed a slight persistence of endo H sensitivity at 1 h of chase compared with KL-1, indicating that passage of these proteins through the Golgi complex was delayed (Fig. 4 A). This is consistent with the delayed maturation of these proteins to their cell surface forms. Most of the KL-Q241 protein remained sensitive to endo H treatment at 1 h of chase, indicating that



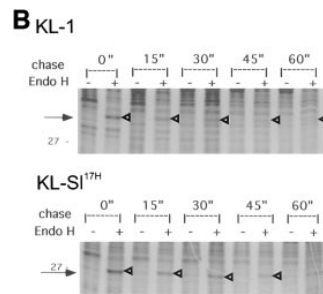
**Figure 3.** Determination of cell surface expression of KL in  $SI^{17H}/SI^{17H}$  fibroblasts by FACS®.

the protein was not properly transiting the Golgi complex. A more thorough time course comparing KL-1 and KL- $SI^{17H}$  showed that at 45 and 60 min of chase, KL- $SI^{17H}$  displayed some persistence of endo H sensitivity, whereas KL-1 had become fully resistant (Fig. 4 B). Thus, KL- $SI^{17H}$  is delayed in exiting the ER. Immunofluorescence staining of KL-1 expressing COS-1 cells revealed a pattern characteristic of a membrane protein which outlined the shape of the cytoplasm (Fig. 5). The KL- $SI^{17H}$  staining pattern was indistinguishable from that of KL-1. KL-Q241 staining was observed at the cell surface as well, but appeared less intense than KL-1 staining. Upon permeabilization, staining patterns of all KL-expressing cells included a reticular pattern surrounding the nucleus. Relative to cell surface staining, KL-Q241 immunofluorescence was more pronounced in this juxtannuclear region, and this intracellular staining pattern appears to overlap with that of an E.R. marker antibody. Therefore, these results are consistent with the idea that the delay in maturation of the KL-Q241 protein and to a lesser extent of the  $SI^{17H}$  protein is a consequence of a delay in passage through a pre-Golgi compartment.

**Efficient Formation of Homodimers of Cell Membrane-associated KL and Cytoplasmic Domain Deletion Variants, and Diminished Formation of Dimers of the  $SI^{17H}$  Protein.** Growth factors and cytokines commonly activate their receptors by mediating receptor dimerization. Recent studies (see references 50 and 51) with the Kit receptor suggest that bivalent binding of KL dimers is driving Kit receptor dimerization. Soluble KL is predominantly in a dimeric state. However, studies of KL dimerization kinetics imply that in vivo, 90% of KL circulating in serum exists as monomers. It is not known if the membrane-associated KL protein on the cell surface is monomeric or dimeric. To address this question, we set out to identify KL dimers by using chemical cross-linking methodology. COS-1 cells transiently transfected to express KL-1 and KL-2 were treated with a hydrophilic and a hydrophobic cross-linker (DSS and BS<sup>3</sup>), cell lysates were

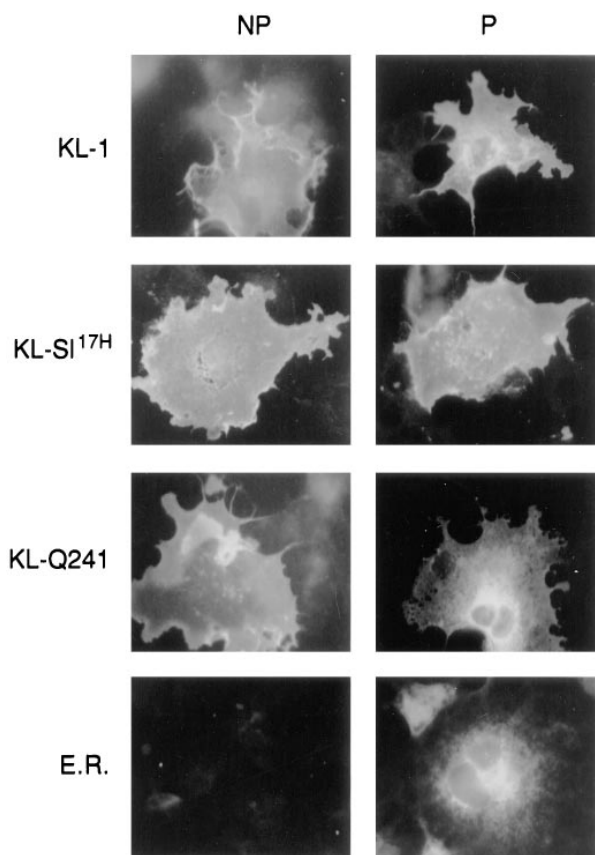


**Figure 4.** Endo H treatment of KL-1, KL cytoplasmic domain variants, and KL- $SI^{17H}$ . COS-1 cells were transfected with different KL constructs and labeled with <sup>35</sup>S-methionine. Cell lysates collected at different time points were immunoprecipitated and treated in the presence or absence of endo H. The reaction products were analyzed by SDS-PAGE (12%). Arrowheads, Immature KL protein products.



then fractionated by SDS-PAGE, and KL protein products were identified by Western blotting (Fig. 6). Both cross-linkers efficiently cross-linked the normal KL-1 and KL-2 transmembrane proteins to produce dimeric molecules of 80–85 kD (Fig. 6). These results suggest that a major fraction of the cell-associated KL-1 and KL-2 molecules are in a dimeric form. It was next of interest to determine whether the cytoplasmic domain sequences of KL and/or the  $SI^{17H}$  cytoplasmic domain sequences affect the formation of cell-associated KL dimers. Whereas deletion of cytoplasmic domain sequences did not affect KL dimer formation as determined by cross-linking (Fig. 6), importantly,  $SI^{17H}$  dimer formation was diminished significantly (Fig. 7), and removal of the  $SI^{17H}$  cytoplasmic domain sequences by deletion restored the ability to form dimers (Fig. 7).

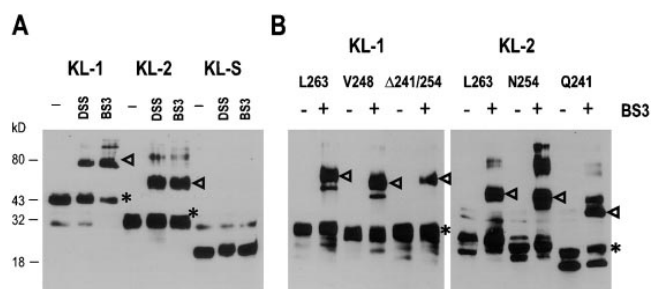
**Cell Adhesion Properties of KL Cytoplasmic Domain Variants.** In an earlier study, Cheng and Flanagan found a significant reduction in binding of a Kit ectodomain alkaline phosphatase fusion protein to COS cells expressing the KL  $SI^{17H}$  protein compared with COS cells expressing the normal KL protein (26). The reduction in KL  $SI^{17H}$  dimerization we observed in COS cells may contribute to the reduced binding of Kit ectodomain. KL/Kit interactions may also be important in cell–cell adhesion. Therefore, we explored the functional consequences of the cytoplasmic domain deletion and  $SI^{17H}$  mutations on cell–cell adhesion.



**Figure 5.** KL protein expression detected by immunofluorescence in COS-1 cells expressing wild-type and mutant KL protein products. COS-1 cells transfected with KL-1, KL-Q241, and KL-*SI*<sup>17H</sup> grown on coverslips were fixed with 2% formaldehyde (NP, nonpermeabilized control) and permeabilized with 0.1% saponin (P). The cells were incubated with anti-KL antibody or an ER marker antibody and FITC-conjugated goat anti-rabbit antibody. Photographs were taken with fluorescent microscopy.

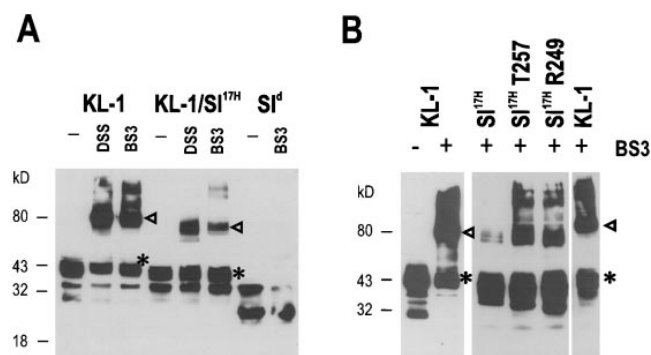
Cell adhesion between kit receptor-expressing mast cells and KL-expressing COS-1 cells or fibroblasts had been demonstrated previously (22, 23). In 24-well dishes, BMMCs ( $1.5 \times 10^5$  cells per well) were plated on top of COS-1-expressing KL-1, KL-N254, KL- $\Delta$ 241/254, KL-Q241, and KL-*SI*<sup>17H</sup>. 72 h after transfection,  $1.5 \times 10^5$  BMMCs were plated per well. After an incubation period of 1 h at 37°C, nonattached BMMCs were removed, and adherent BMMCs were counted by visual inspection in a microscope. COS cells expressing KL-1, KL-N254, and KL- $\Delta$ 241/254 had equivalent numbers of BMMCs attached to the cell surface, and attachment was inhibited completely by the antagonistic anti-*c-kit* mAb ACK2 (38) (Fig. 8). Adhesion of BMMCs to COS-1 cells expressing KL-Q241 was reduced dramatically (Fig. 8). Similar numbers of BMMCs attached to COS-1 cells transfected with either KL-Q241, the secretory KL-1S (2), or a control plasmid. Interestingly, the number of BMMCs attached to COS cells expressing KL-*SI*<sup>17H</sup> was only about half of that attached to COS cells expressing KL-1 (Fig. 8).

**Effects of the *SI*<sup>17H</sup> Mutation on Hematopoiesis In Vivo and In Vitro.** Previous analysis of mutant *SI*<sup>17H</sup>/*SI*<sup>17H</sup> mice sug-

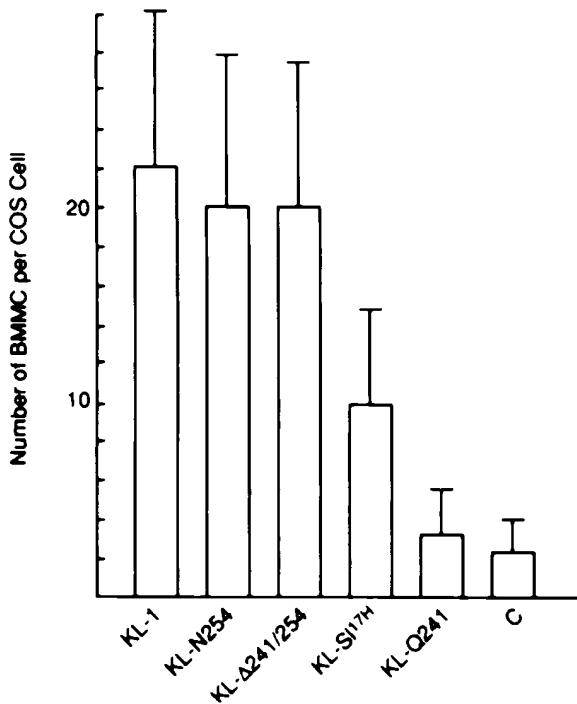


**Figure 6.** Demonstration of KL-1, KL-2 dimers in transfected COS-1 cells. COS-1 cells were transfected with various normal and mutant KL-1 and KL-2 constructs: (A) KL-1, KL-2, and KL-S; (B) KL-1: L263, V248,  $\Delta$ 241/254; KL-2: L263, N254, Q241. 72 h after transfection, they were treated with the cross-linkers DSS and BS<sup>3</sup> for 1 h at 37°C. Cell lysates were collected and fractionated by SDS-PAGE (12%), and KL protein products were identified by Western blotting. Arrowheads, KL dimers. \*Monomers. Molecular mass (M<sub>r</sub>) markers are indicated in kilodaltons.

gested mild effects in the hematopoietic system, i.e., a mild macrocytic anemia was observed (29). We have reexamined the effects of the *SI*<sup>17H</sup> mutation on the hematopoietic system in vivo as well as in vitro. Careful analysis of the in vivo hematopoietic parameters indicates that hematocrit values, WBC, neutrophil, and platelet numbers, as well as bone marrow cellularity are normal in homozygous mutant mice (Table 1). In contrast, the numbers of skin mast cells were reduced by 45% and peritoneal mast cells by 85%, indicating an effect of the *SI*<sup>17H</sup> mutation on mast cell development and/or number (Table 1). In agreement with a lack of an erythroid deficiency, in vitro analysis of hematopoietic progenitors revealed normal numbers of BFU-E. However, mixed colonies (CFU-GEMM) and CFU-GM were reduced slightly, by  $\approx$ 33%. We also established long term bone marrow cultures and followed output of he-



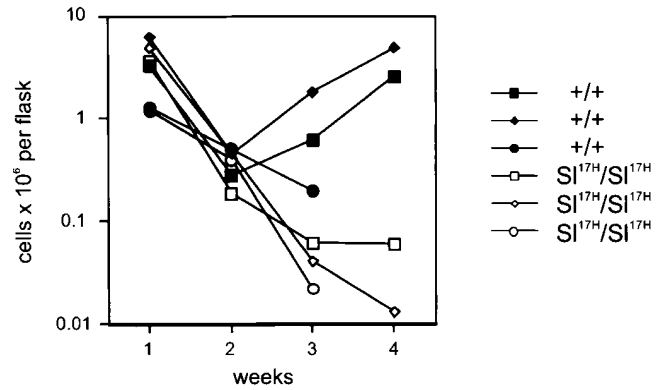
**Figure 7.** The *SI*<sup>17H</sup> mutation affects dimer formation on the cell surface. COS-1 cells were transfected with normal KL-1 constructs and mutant *SI*<sup>17H</sup> constructs containing various COOH-terminal deletions: *SI*<sup>17H</sup>-T257, *SI*<sup>17H</sup>-R249, and *SI*<sup>17H</sup>-A240. 72 h after transfection, cell surface proteins were biotinylated, and cells were treated with the cross-linkers DSS and BS<sup>3</sup> (as indicated) for 1 h at 37°C. Cell lysates were fractionated on streptavidin-sepharose columns, the biotinylated fraction was electrophoretically fractionated by SDS-PAGE (12%), and KL protein products were identified by Western blotting. Arrowheads, KL dimers. \*Monomers. Molecular mass (M<sub>r</sub>) markers are indicated in kilodaltons.



**Figure 8.** Adhesion of BMMCs to COS-1 cells expressing KL-1, secretory KL-1S, different KL cytoplasmic variants, and KL- $S^{I^{7H}}$ . COS-1 cells were transfected to 24-well plates 24 h after transfection at a density of  $7.5 \times 10^4$  cells per well. BMMCs ( $1.5 \times 10^5$  cells per well) were incubated with transfected COS-1 cells for 1 h in the presence and absence of anti-c-kit antibody. Nonattached BMMCs were washed away three times with medium, and the number of attached BMMCs was counted by microscopic inspection. The number of BMMCs attached to individual COS cells is shown.

matopoietic cells at weekly intervals. These experiments showed that cultures from  $S^{I^{7H}}/S^{I^{7H}}$  bone marrow were not able to sustain long-term production of hematopoietic cells (Fig. 9). This result is reminiscent of the defect observed in cultures derived from bone marrow of  $SI/SI^d$  mice, which produce only the soluble form of KL and no membrane-associated KL.

Kit-mediated adhesion of hematopoietic cells is known to be mediated by at least two distinct mechanisms: (a) tethering via the membrane growth factor-receptor inter-



**Figure 9.** Long-term culture of bone marrow cells from  $S^{I^{7H}}/S^{I^{7H}}$  mice. Nonadherent cells were counted at weekly intervals. Results from three separate experiments are shown.

action, and/or (b) by inside-out activation of integrin-mediated adhesion (very late antigen [VLA]-4 and VLA-5) (22, 23, 39, 40). Furthermore, it has been established that VLA-4-mediated adhesion is critical for homing or lodging of early hematopoietic progenitors to the bone marrow in transplantation experiments (41, 42). Therefore, we determined whether  $S^{I^{7H}}/S^{I^{7H}}$  mice were suitable recipients in spleen colony assays using normal donor bone marrow. Interestingly, we observed a reduced number of spleen colonies (65% of normal) in  $S^{I^{7H}}/S^{I^{7H}}$  recipient mice (Table 2). This may suggest that the  $S^{I^{7H}}$  mutation impairs lodging/homing of transplanted CFU-S progenitors to the spleen of irradiated mice.

## Discussion

The  $S^{I^d}$  allele arose as a result of an intragenic deletion including the transmembrane domain and COOH terminus, generating a secreted KL protein product with normal biological activity (2, 22, 27). Analysis of the  $S^{I^d}$  phenotype has been of great value in understanding the differential biological roles of membrane-associated and soluble forms of KL. The biological characteristics of homozygous  $S^{I^d}/S^{I^d}$  mice and of  $SI/SI^d$  mice indicate that the  $S^{I^d}$  protein supports some level of KL function. However,  $S^{I^d}$  mice display major defects in facilitating proliferation and survival of tar-

**Table 1.** Mast Cells and Hematopoietic Progenitors in  $S^{I^{7H}}/S^{I^{7H}}$  and C3H Control Mice

Genotype	Skin mast cells	Peritoneal mast cells	Bone marrow cellularity	CFU-GEMM	CFU-GM	BFU-E
$+/+$						
Exp. 1	162.5 ± 4.5	49.5 ± 9.9	11.54 ± 0.77	2,626 ± 335	29,003 ± 2,871	1,955 ± 430
Exp. 2				770 ± 191	9,717 ± 1,215	2,677 ± 204
$S^{I^{7H}}/S^{I^{7H}}$						
Exp. 1	91.6 ± 12.8	8.0 ± 1.6	12.42 ± 1.67	1,634 ± 608	19,923 ± 3,163	1,925 ± 515
Exp. 2				583 ± 210	10,266 ± 1,811	2,391 ± 657

**Table 2.** Reciprocal Transplantation of Bone Marrow from  $Sl^{l7H}/Sl^{l7H}$  and Control C3H Mice into Irradiated Control and  $Sl^{l7H}/Sl^{l7H}$  Mice

Donor	Recipient	CFU-S/ $10^5$ cells
+/+	+/+	25.3 ± 2.1
$Sl^{l7H}/Sl^{l7H}$	+/+	36.9 ± 2.0 $P = 0.0008$
+/+	+/+	25.1 ± 1.7
+/+	$Sl^{l7H}/Sl^{l7H}$	16.5 ± 1.0 $P = 0.0024$

get cells. Therefore, the cell-associated form of KL plays a critical role in *c-kit* function, and the cytoplasmic domain of KL is potentially important to the processes mediated by juxtacrine signaling. This notion is supported by the mutant phenotypes of the  $Sl^{l7H}$  allele, a splice site mutation that results in the substitution of amino acids 239–273 in the KL cytoplasmic domain with 27 extraneous amino acids (28). Therefore, the  $Sl^{l7H}$  allele provided an opportunity to analyze the in vitro and in vivo consequences of cytoplasmic domain modification.

The cytoplasmic domain of KL is highly conserved in evolution (Fig. 1), yet very little is known about its function. We have attempted to elucidate the roles of KL cytoplasmic domain sequences as they relate to biosynthetic processing, cell adhesion, and juxtacrine signaling by using in vitro and in vivo genetic approaches. The major conclusions of our study are (a) that cytoplasmic domain sequences are important for biosynthetic processing of KL through the ER and Golgi complex and to the cell surface; (b) that the membrane forms of KL exist as homodimers on the cell surface and that dimerization may be an essential step in KL/Kit-mediated juxtacrine signaling; and (c) that analysis of in vivo phenotypes of  $Sl^{l7H}/Sl^{l7H}$  mice revealed Kit-dependent processes in hematopoiesis in which membrane KL is limiting, and they suggest a role for Kit in homing of hematopoietic progenitors to spleen.

Our findings that the KL cytoplasmic domain is required for normal processing to the cell surface are consistent with reports on a variety of secreted or membrane-anchored proteins in which cytoplasmic domain mutations disrupted intracellular trafficking and maturation. For example, mutations removing the four COOH-terminal cytoplasmic residues of  $\alpha$ -1 proteinase inhibitor or the COOH-terminal 22 amino acids of thyroxine-binding globulin caused nascent protein to be retained in the ER with resultant lack of secretion (43, 44). Additionally, single point mutations of glycines in the cytoplasmic domain of *P*-glycoprotein or a single point mutation in the cytoplasmic kinase domain of the Kit receptor caused these proteins which are normally membrane-anchored to be inefficiently glycosylated and retained in the ER (45, 46). Cytoplasmic domain mutant proteins may not interact properly with molecules involved with transport from the ER to the cell surface. Also, incor-

rect folding as a consequence of cytoplasmic domain mutations may be responsible for the maturation defects (47).

Soluble KL forms noncovalent dimers in solution (15, 48, 49), and it is thought that dimerization of the ligand as well as of the receptor are essential steps for receptor activation. Therefore, it seemed reasonable that juxtacrine signaling by cell-associated forms of KL may involve membrane KL dimers. Our finding of efficient dimer formation of cell-associated KL-1 and KL-2 molecules on transfected COS cells supports this conjecture. Importantly, the apparent large fraction of KL dimers on the cell surface in the absence of cognate receptor implies that dimer formation may not be a rate-limiting event in juxtacrine signaling between KL and Kit. This is in contrast to receptor tyrosine kinases such as Kit which primarily exist as monomers on the cell surface and which form dimers in response to engagement with the cognate ligand (50, 51). Our analysis of various cytoplasmic domain deletion mutations suggests that the cytoplasmic domain of KL does not have an active role in dimer formation. The decreased efficiency of dimer formation of the  $Sl^{l7H}$  protein suggests that the nonsense cytoplasmic domain sequence of  $Sl^{l7H}$  prevents normal KL dimerization (possibly by steric hindrance), and this may contribute to the phenotypic defects of  $Sl^{l7H}$  mice.

Cell adhesion assays indicated that Kit-mediated attachment of BMSCs to COS cells expressing KL- $Sl^{l7H}$  was  $\approx 50\%$  of the KL-1 control. This was in contrast to KL-N254 and KL- $\Delta 241$ -254, where no significant reduction in adhesion was seen. Interestingly, while a similar delay in the processing of  $Sl^{l7H}$ , KL-N254, and KL- $\Delta 241$ -254 to the cell surface is observed,  $Sl^{l7H}$  specifically displays reduced cell surface dimerization. This may suggest that the adhesion defect of  $Sl^{l7H}$  is a consequence of reduced  $Sl^{l7H}$  cell surface dimer formation. Therefore, the KL- $Sl^{l7H}$  protein is expressed on the cell surface, but the abnormal cytoplasmic domain sequences appeared to interfere with membrane KL-mediated adhesion of BMSCs and juxtacrine signaling.

The  $Sl^{l7H}$  mutation affects melanogenesis and gametogenesis (28, 29). A mild effect on hematopoiesis of  $Sl^{l7H}$  has also been reported. We have now investigated the hematopoietic parameters in  $Sl^{l7H}/Sl^{l7H}$  mice in detail. Although bone marrow cellularity, hematocrit, and WBC, neutrophil, and platelet counts appeared to be normal, the number of tissue mast cells was appreciably affected by the  $Sl^{l7H}$  mutation. In addition, hematopoietic progenitor numbers in the bone marrow of  $Sl^{l7H}/Sl^{l7H}$  mice did not deviate significantly from normal controls, although slightly reduced numbers of mixed colonies (CFU-GEMM) and slightly increased numbers of CFU-S were observed. Evidence that the hematopoietic microenvironment is compromised in  $Sl^{l7H}/Sl^{l7H}$  mice comes from two different experiments. First, reciprocal transplantation of normal bone marrow into  $Sl^{l7H}/Sl^{l7H}$  and normal mice revealed a reduced number of spleen colonies in the  $Sl^{l7H}/Sl^{l7H}$  recipients, suggesting that homing or lodging to the spleen is affected by the mutation. Second, the bone marrow from  $Sl^{l7H}/Sl^{l7H}$  mice



does not support long-term culture of hematopoietic progenitors. The biochemical defects we observed for the  $Sl^{7H}$  protein, including delayed processing and reduced dimer formation, may contribute to these hematopoietic deficiencies. The observation that homing or lodging of progenitors to the spleen of irradiated mice is affected is of great interest. KL had been shown to mediate adhesion processes in hematopoietic cells and mast cells by means of an inside-out activation of the integrins VLA-4 and VLA-5 (39, 40, 52, 53). Also, VLA-4-mediated adhesion has been demonstrated to play a role in homing or lodging of hematopoietic progenitors to the spleen (41). In addition, tethering via the membrane growth factor-receptor interaction is thought to provide a mechanism for KL/Kit-mediated cell-cell adhesion (2, 22). Although our results provide evidence that supports a role for KL/Kit in homing or lodging of hematopoietic progenitors to the spleen possibly by an integrin-mediated mechanism, a role for KL/Kit in homing to the bone marrow as suggested by Papayannopoulou and Craddock (54) remains to be investigated.

The number of germ cells in neonatal gonads is reduced similarly in male and female  $Sl^{7H}/Sl^{7H}$  mice as a result of an effect on primordial germ cells; however, the  $Sl^{7H}$  allele affects male but not female fertility (28, 55). In postnatal spermatogenesis, KL-2 is more abundant than KL-1, whereas

in oogenesis, KL-1 and KL-2 are equally abundant (2, 32). Since the KL-2 protein is more resistant to proteolytic processing and therefore is more stable on the cell surface, the preferred expression of KL-2 in testis development implies a more selective role for transmembrane KL in spermatogenesis. Therefore, the defect in dimerization of  $Sl^{7H}$  would more severely affect spermatogenesis and may in part account for the sterility of  $Sl^{7H}$  males, while females remain fertile. In addition, immunohistochemical studies showed that the transmembrane KL protein accumulates in the basolateral region of the seminal epithelium in Sertoli cells at the time when *c-kit*-expressing spermatogonia begin to mature (32). KL-expressing Sertoli cells supporting KL/*c-kit*-dependent spermatogonial proliferation have the morphology of a polarized epithelium. Possibly, the cytoplasmic domain of KL may direct sorting of this protein to the basolateral region of Sertoli cells and/or affect its stability in the basolateral region of these cells. In the adult ovary, follicle cells surrounding the oocyte are less polarized than Sertoli cells. Therefore, although a 50% decrease in  $Sl^{7H}$  protein may be tolerated in oogenesis, misdirected expression of KL- $Sl^{7H}$  in polarized Sertoli cells may affect spermatogonial proliferation and survival and result in male sterility.

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