

## High Levels of Human $\gamma$ -Globin Gene Expression in Adult Mice Carrying a Transgene of Deletion-Type Hereditary Persistence of Fetal Hemoglobin

MURAT O. ARCASOY,<sup>1</sup> MARC ROMANA,<sup>1†</sup> MARY E. FABRY,<sup>2</sup> EVA SKARPIDI,<sup>3</sup>  
RONALD L. NAGEL,<sup>2</sup> AND BERNARD G. FORGET<sup>1\*</sup>

*Department of Medicine, Yale University School of Medicine, New Haven, Connecticut 06510<sup>1</sup>; Department of Medicine, Albert Einstein College of Medicine, Bronx, New York 10461<sup>2</sup>; and Division of Medical Genetics, University of Washington School of Medicine, Seattle, Washington 98195<sup>3</sup>*

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**Persistent expression of the  $\gamma$ -globin genes in adults with deletion types of hereditary persistence of fetal hemoglobin (HPFH) is thought to be mediated by enhancer-like effects of DNA sequences at the 3' breakpoints of the deletions. A transgenic mouse model of deletion-type HPFH was generated by using a DNA fragment containing both human  $\gamma$ -globin genes and HPFH-2 breakpoint DNA sequences linked to the core sequences of the locus control region (LCR) of the human  $\beta$ -globin gene cluster. Analysis of  $\gamma$ -globin expression in six HPFH transgenic lines demonstrated persistence of  $\gamma$ -globin mRNA and peptides in erythrocytes of adult HPFH transgenic mice. Analysis of the hemoglobin phenotype of adult HPFH transgenic animals by isoelectric focusing showed the presence of hybrid mouse  $\alpha_2$ -human  $\gamma_2$  tetramers as well as human  $\gamma_4$  homotetramers (hemoglobin Bart's). In contrast, correct developmental regulation of the  $\gamma$ -globin genes with essentially absent  $\gamma$ -globin gene expression in adult erythroid cells was observed in two control non-HPFH transgenic lines, consistent with autonomous silencing of normal human  $\gamma$ -globin expression in adult transgenic mice. Interestingly, marked preferential overexpression of the LCR-distal  $\gamma$ -globin gene but not of the LCR-proximal  $\gamma$ -globin gene was observed at all developmental stages in erythroid cells of HPFH-2 transgenic mice. These findings were also associated with the formation of a DNase I-hypersensitive site in the HPFH-2 breakpoint DNA of transgenic murine erythroid cells, as occurs in normal human erythroid cells *in vivo*. These results indicate that breakpoint DNA sequences in deletion-type HPFH-2 can modify the developmentally regulated expression of the  $\gamma$ -globin genes.**

The human  $\beta$ -like globin gene cluster is located on the short arm of chromosome 11 and contains five linked functional globin genes and an upstream locus control region (LCR). The tissue-specific expression of the embryonic, fetal, and adult globin genes is developmentally regulated with sequential activation and silencing of individual genes during ontogeny, characterized by two switches in the pattern of  $\beta$ -like globin gene expression during development. The first switch is observed early in development as the major site of hematopoiesis shifts from the embryonic yolk sac to the fetal liver and is characterized by the high-level expression of the fetal  $\gamma$ - and  $\epsilon$ -globin genes that replaces the expression of the embryonic  $\epsilon$ -globin gene. A second switch, from  $\gamma$ - to  $\delta$ - and  $\beta$ -globin gene expression, occurs late in gestation as well as in the perinatal period and results in high-level expression of the adult  $\beta$ -globin genes in the erythroid cells from the bone marrow, replacing the expression of the fetal  $\gamma$  genes following birth (6, 9, 39, 44). The exact mechanisms that regulate globin gene switching are as yet incompletely understood. However, developmental regulation of this multigene locus appears to be mediated by complex developmental stage- and tissue-specific interactions

between *cis*- and *trans*-acting regulatory elements within the gene cluster.

Genetic disorders that alter hemoglobin (Hb) switching have provided naturally occurring molecular models for the study of the regulation of globin gene transcription and the mechanisms of Hb switching during development. Naturally occurring deletions within the  $\beta$ -globin cluster that interfere with fetal-to-adult Hb switching result in two related but discrete clinical syndromes,  $\delta\beta$ -thalassemia and hereditary persistence of fetal Hb (HPFH), which are characterized by inappropriate, persistent expression of the fetal  $\gamma$ -globin genes in adult life. Individuals heterozygous for deletion-type HPFH have normal erythrocyte (RBC) parameters, but their Hb consists of 20 to 30% Hb F which is distributed in a pancellular fashion among the RBCs. In contrast, individuals with heterozygous  $\delta\beta$ -thalassemia have smaller-than-normal RBCs and lower Hb F levels (10 to 15%), distributed in a heterocellular fashion (5, 34, 39). One hypothesis to explain persistent  $\gamma$ -globin gene expression in adult life in these conditions is that the deletions remove sequences downstream of the  $\gamma$ -globin genes which are necessary for the silencing of the genes in adult life (25). Another mechanism for persistent high levels of Hb F expression could involve the introduction into the vicinity of the  $\gamma$ -globin genes of DNA sequences from the breakpoint regions that can influence  $\gamma$ -globin gene expression following birth through enhancer-like effects (1, 18, 35, 42). These two mechanisms are not mutually exclusive, however, and a different balance of these and possibly other effects could play a role in the different phenotypes of persistent  $\gamma$ -globin gene expression observed in adults with HPFH and  $\delta\beta$ -thalassemia.

\* Corresponding author. Mailing address: Yale University School of Medicine, Hematology Section, WWW 403, 333 Cedar St., New Haven, CT 06510. Phone: (203) 785-4144. Fax: (203) 785-7232. E-mail: bernard.forget@yale.edu.

† Present address: Unité de Recherche sur la Drepanocytose, INSERM U-359, Centre Hospitalier Universitaire, Guadeloupe, French West Indies.

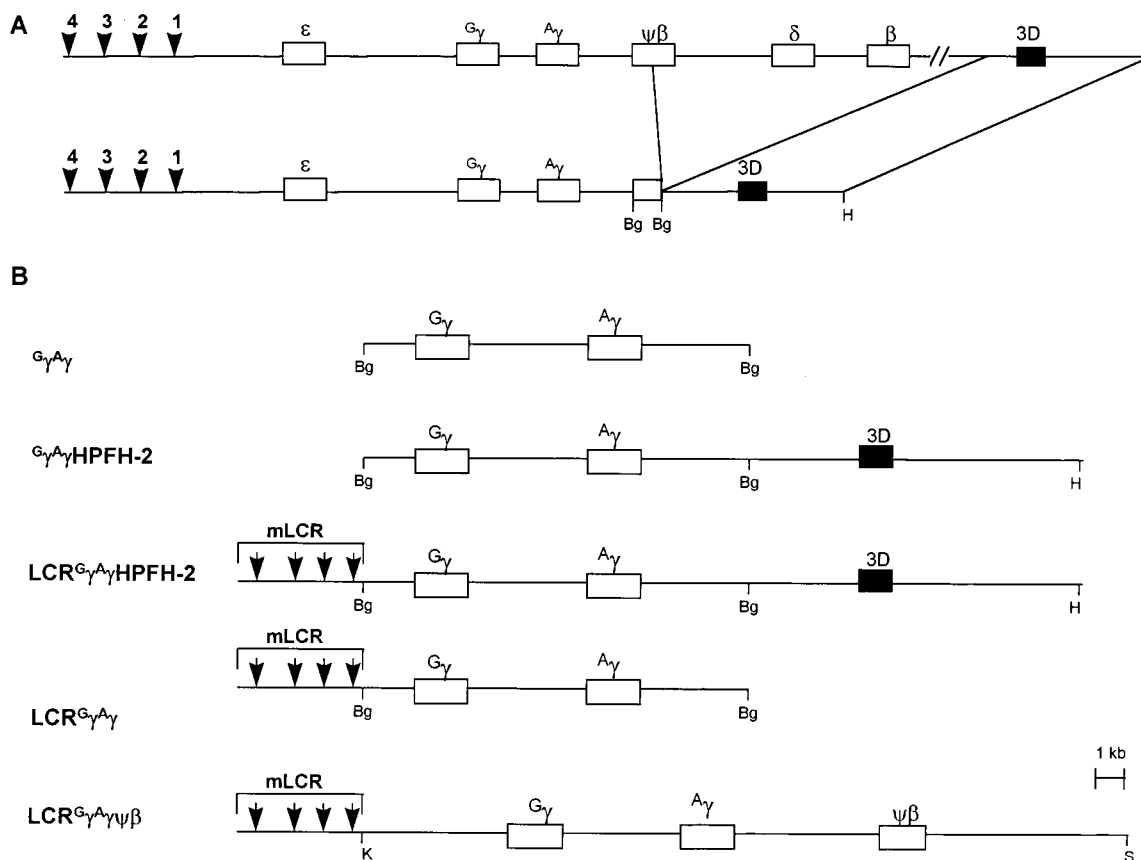


FIG. 1. (A) Diagrams of the normal and mutant (HPFH-2) human  $\beta$ -globin loci. The arrowheads indicate the DNase I-HSSs in the LCR. (B) DNA fragments that were used to generate transgenic mice. See Materials and Methods for a detailed description of the DNA fragments used in microinjection. mLCR, mini-LCR cassette; Bg, *Bgl*I; H, *Hind*III; K, *Kpn*I; S, *Sal*I.

A large body of experimental results has been generated by the analysis of a large number of different transgenic mice carrying a variety of fragments from the human  $\beta$ -globin gene cluster. In the absence of the LCR, individual human  $\gamma$ -globin transgenes with short segments of flanking DNA have been shown to be expressed during mouse development in yolk sac-derived but not adult erythroid cells (7, 26). These early results suggested that DNA sequences necessary to mediate developmental stage-specific expression of the  $\gamma$ -globin genes are located within or near the genes themselves. However, later experiments by Enver et al. (16) and Behringer et al. (3) showed that when linked with the LCR,  $\gamma$ -globin transgenes in the absence of linked  $\beta$ -globin genes were expressed in both embryonic and adult RBCs and that silencing of  $\gamma$  genes in adult erythroid cells was achieved by linking the LCR to the intact  $\gamma\delta\beta$  region. These results suggested a competitive model for regulation of the  $\gamma$ - and  $\beta$ -globin genes in transgenic mice (12, 15, 16, 23, 24). Subsequent experiments by Dillon and Grosfeld (11) showed that  $\gamma$ -globin genes linked to the LCR are in fact silenced autonomously in adult erythroid tissues in the absence of an adjacent competing  $\beta$ -globin gene. Experiments with transgenic mice carrying a 70-kb transgene containing the entire human  $\beta$ -globin locus, including its LCR, in their normal structural configuration, as well as analyses of mice carrying the total  $\beta$ -globin gene cluster and LCR in large yeast artificial chromosome (YAC) transgenes, showed that the human  $\gamma$ -globin genes are correctly regulated during development and are expressed during the embryonic and early fetal

stages of murine erythropoiesis (20, 32, 41). The phenotype of mice carrying large transgenes of the  $\beta$ -globin gene cluster but with deletions of both the  $\delta$ - and  $\beta$ -globin genes demonstrated autonomous silencing of the  $\gamma$ -globin genes in adult transgenic mice, supporting the possible role of 3' breakpoint DNA sequences in the generation of  $\delta\beta$ -thalassemia and deletion-type HPFH phenotypes (33, 41).

To investigate the molecular mechanisms of  $\gamma$ -globin gene overexpression in adults with deletion-type HPFH, we have studied a model for the two most common forms of deletion-type  $G\gamma^A\gamma$  pancellular HPFH, HPFH-1 and HPFH-2. These disorders are associated with extensive deletions of the  $\beta$ -globin cluster involving approximately 105 kb of DNA (10) that includes both the  $\delta$ - and  $\beta$ -globin genes (Fig. 1A). The purpose of this study was to test the functional significance of the breakpoint DNA sequences in deletion-type HPFH and to determine whether the presence of the  $\beta$ -cluster LCR is required for generation of the HPFH phenotype in transgenic mice carrying a 25-kb human DNA fragment mimicking the structure of the  $\beta$  cluster in HPFH-2. We generated two transgenic lines carrying the HPFH-2 transgene without the LCR and six transgenic lines with the HPFH-2 transgene with the LCR. Analysis of the phenotypes of these transgenic lines indicates that adult mice carrying the HPFH-2 transgene without the LCR displayed a delay in the silencing of  $\gamma$ -gene expression during fetal development but did not have persistent  $\gamma$ -gene expression in adult erythroid cells. On the other hand, analysis of the HPFH-2 transgenic lines with the LCR revealed

that human  $\gamma$ -globin mRNA and peptide chains were persistently expressed at significant levels and incorporated into mouse  $\alpha_2$ -human  $\gamma_2$  hybrid Hb tetramers. This persistent  $\gamma$ -gene expression was associated with formation of a DNase I-hypersensitive site (HSS) within HPFH-2 3' breakpoint DNA region 3D in erythroid tissues of adult mice with the LCR/HPFH-2 transgene. In contrast, correct developmental regulation of  $\gamma$ -globin gene expression was observed in animals from two different control non-HPFH transgenic lines carrying LCR<sup>G $\gamma$</sup>  and LCR<sup>G $\gamma$</sup>  $\psi\beta$  transgenes in which there was appropriate silencing of the  $\gamma$ -globin transgenes in adult erythroid cells.

#### MATERIALS AND METHODS

**DNA fragments.** The following human DNA fragments shown in Fig. 1B were prepared for injection into mouse oocytes. (i) G $\gamma$  is a 13-kb *Bgl*II fragment containing both  $\gamma$ -globin genes and downstream sequences up to approximately the 5' HPFH-2 breakpoint site. (ii) G $\gamma$ HPFH-2 is a 25-kb transgene containing 12 kb of HPFH-2 3' breakpoint DNA sequences linked to G $\gamma$ . (iii) LCR<sup>G $\gamma$</sup> HPFH-2 is an ~30-kb transgene containing a 4.7-kb LCR cassette linked to the 25-kb fragment. (iv) LCR<sup>G $\gamma$</sup>  is an ~18-kb transgene with the LCR cassette linked to the 13-kb *Bgl*II genomic DNA fragment. (v) LCR<sup>G $\gamma$</sup>  $\psi\beta$  is an ~32-kb transgene containing the LCR cassette, both  $\gamma$ -globin genes, and 14 kb of downstream sequences ending 5' to the  $\delta$ -globin gene. The 13-kb *Bgl*II genomic DNA fragment containing both the G $\gamma$  and A $\gamma$  genes (GenBank Humhbb coordinates: 32820 to 45698) was purified from a cosmid clone containing the  $\beta^S$ -globin cluster described previously (8) and subcloned into the *Bam*HI site of cosmid vector pWE15, which contained a modified polylinker. To construct the LCR<sup>G $\gamma$</sup> HPFH-2 transgene, a 12-kb *Bgl*III-*Hind*III genomic DNA fragment from a  $\lambda$  phage clone ( $\lambda$ B-1) containing the normal DNA overlapping the 3' breakpoint of HPFH-1 and HPFH-2 (42) was subcloned downstream of the 13-kb *Bgl*III genomic fragment containing the  $\gamma$ -globin genes to yield a 25-kb DNA fragment whose structure is virtually identical to the structure of HPFH-2 DNA.

An LCR cassette was constructed by ligating 5' HSS1 and HSS2 isolated as a 1.4-kb *Pst*I-*Hind*III fragment from the  $\mu$ LAR plasmid (19) to a 1.2-kb *Xmn*I-*Hind*III fragment containing HSS3 and a 2.0-kb *Bam*HI-*Xba*I fragment containing HSS4 (GenBank Humhbb coordinates: HSS1, 13062 to 13769; HSS2, 8486 to 9218; HSS3, 3975 to 5172; HSS4, 308 to 2352). This 4.7-kb LCR cassette was subcloned in the genomic orientation upstream of the  $\gamma$ -globin genes in the G $\gamma$ HPFH-2 DNA fragment to yield the LCR<sup>G $\gamma$</sup> HPFH-2 transgene. The first control transgene, LCR<sup>G $\gamma$</sup> , contained the LCR cassette, which was subcloned in the genomic orientation upstream of the 13-kb *Bgl*III genomic fragment containing the  $\gamma$ -globin genes up to the HPFH-2 5' breakpoint. The second control transgene, LCR<sup>G $\gamma$</sup>  $\psi\beta$ , was obtained by inserting the LCR cassette, in the genomic orientation, upstream of a 27-kb *Kpn*I-*Sal*I genomic fragment (GenBank Humhbb coordinates: 27250 to 54726) containing the G $\gamma$ - and A $\gamma$ -globin genes and downstream sequences including the  $\psi\beta$  pseudogene up to the  $\delta$ -globin gene.

**Transgenic mouse lines.** The DNA fragments were prepared for microinjection as described previously (40). A solution containing 10 ng of DNA per  $\mu$ l was injected into the pronuclei of (C57BL/6  $\times$  SJL)<sub>F2</sub> zygotes and transferred to pseudopregnant females. Transgenic mice were identified as described by Starck et al. (40), and transgenic founder mice were bred with nontransgenic B6/SJL mates to establish transgenic lines. Studies of transgene expression during development were performed with each transgenic line by mating hemizygous transgenic males with nontransgenic B6/SJL females to obtain embryos from timed pregnancies at 11.5, 13.5, and 16.5 days postcoitum. The morning on which the mating plug was observed was designated day 0.5. The copy numbers of the transgenes were determined by digestion of 10  $\mu$ g of genomic DNA from the tail skin of progeny of two or three different animals from each line. Digestion with *Eco*RI, *Bam*HI, or *Sac*I was carried out and followed by 0.8% agarose gel electrophoresis and Southern blotting as described by Sambrook et al. (36). Several different probes from the transgenes were hybridized to the blots, and the hybridization signals were quantitated with a PhosphorImager (Molecular Dynamics). The measurements were made several times, and averages were taken. Copy numbers were determined by comparing the signals from the human transgene to signals from mouse GATA-1 as described by Starck et al. (40).

**RNA analysis.** Blood samples were obtained from mouse embryos and adult mice as described previously (40), and total RNA was isolated by using TRIzol reagent (Gibco-BRL) according to the manufacturer's protocol. Human and murine globin mRNAs were analyzed by quantitative RNase protection assays. The human  $\gamma$ -globin probe was a 403-bp *Nae*I-*Bam*HI fragment (IVS1 deleted) in pGEM4 (Promega) containing 5' upstream sequences and exons 1 and 2 of the  $\gamma$ -globin genes. The antisense probe was synthesized by transcription with Sp6 polymerase to identify a 350-bp protected fragment. For the differential detection of G $\gamma$ - and A $\gamma$ -globin transcripts, the G $\gamma$  probe was transcribed with T7 polymerase from pBluescript (Stratagene) containing a 721-bp *Hinc*II-*Hind*III

G $\gamma$ -globin fragment and the A $\gamma$  probe was transcribed with Sp6 polymerase from pBluescript containing a 980-bp *Pvu*II A $\gamma$ -globin fragment, both of which contained the third exon including the 3' untranslated regions of the G $\gamma$ - and A $\gamma$ -globin genes, respectively, where four consecutive base differences are present between the G $\gamma$  and A $\gamma$  sequences beginning 3 bases 3' to the termination codon. Both probes give a 215-bp protected fragment with the respective G $\gamma$  or A $\gamma$  transcript. Different riboprobes were used for the differential detection of G $\gamma$ - and A $\gamma$ -globin transcripts in the experiments with the mouse lines carrying the transgenes without the LCR. These probes and the riboprobes for detection of murine  $\alpha$ -globin and  $\zeta$ -globin mRNAs and quantitative mRNA analysis methods used for the mouse lines carrying the transgenes without the LCR have been described previously (40). For the mRNA analysis of samples from transgenic mouse lines with the LCR, three different probes (human  $\gamma$ -globin and mouse  $\alpha$ - and  $\zeta$ -globins) of  $1.5 \times 10^6$  cpm total were simultaneously hybridized to 500 ng of total RNA from 11.5-day RBCs and 100 to 200 ng of RNA from RBCs of 13.5- and 16.5-day fetuses as well as newborn and adult mice. Samples were digested with RNases A (2.5 U/ml) and T<sub>1</sub> (100 U/ml). Conditions of probe excess were confirmed in separate experiments. The protected fragments were detected by autoradiography after electrophoresis in 8% polyacrylamide-8-mol/liter urea gels. Quantitation of human and murine mRNAs was performed with a PhosphorImager.

**Isoelectric focusing (IEF).** Hb samples were separated on IEF by using a Resolve-Hb kit (Isolab, Akron, Ohio). Hb concentrations were adjusted to between 2 and 4.5 g% by using the supplied sample preparation solution containing 0.05% KCN. Gels were run until sharp bands formed, usually 2 h. To determine the percentage of each band on the IEF gel, the gel was fixed with 10% trichloroacetic acid solution; soaked in distilled water for 15 min; stained with the JB-2 Staining System (Isolab) containing *o*-dianisidine, a heme-specific stain; and scanned at 520 nm with a model 710 densitometer (Corning, Medfield, Mass.). To identify bands, the band was removed from the unfixed gel, eluted with distilled water, spun to remove gel, and analyzed by high-performance liquid chromatography (HPLC).

**HPLC.** The globin chain composition of mouse RBCs was determined by HPLC using a denaturing solvent that separates the globin chains and a Vydac large-pore (300-Å) C<sub>4</sub> column (4.6 by 250 mm; Separations Group, Hesperia, Calif.) with a modified acetonitrile-H<sub>2</sub>O-trifluoroacetic acid (TFA) gradient similar to that used by Schroeder et al. (37) for separating human globin chains. Two buffers were used: A (0.18% TFA in 36% acetonitrile) and B (0.18% TFA in 46% acetonitrile). Starting with 35% B, the amount of buffer B was increased by 0.67%/min until all of the globin chains were eluted.

**Determination of oxygen affinity of hemoglobin (P<sub>50</sub>).** Blood samples were collected from the mouse tail and placed directly into heparinized mouse saline (330 mosM). The samples were washed three times with 10 mM HEPES buffer containing 5 mM KCl, 5 mM glucose, and enough NaCl to adjust the osmolality to 330 mosM. The buffer pH was 7.4 at 37°C. O<sub>2</sub> equilibrium curves were obtained by running the samples at 37°C on a Hem-O-Scan O<sub>2</sub> Dissociation Analyzer (Aminoco, Silver Spring, Md.) at hematocrits between 10 and 20%.

**DNase I hypersensitivity assays.** Adult mice carrying the LCR<sup>G $\gamma$</sup> HPFH-2 transgene were made anemic by three injections of 0.2 ml of 0.4% acetyl phenylhydrazine per 25 g of body weight and sacrificed 5 days after the start of the injections to harvest the spleen. The spleen was placed in a 10-ml syringe and passed through an 18-gauge hypodermic needle several times and resuspended in RSB buffer (10 mM Tris · HCl [pH 7.4], 10 mM NaCl, 3 mM MgCl<sub>2</sub>). Intact nuclei were isolated as described elsewhere (43), and aliquots were subjected to DNase I digestion at various concentrations. The genomic DNA was purified by standard procedures (36). Appropriate restriction endonucleases were used to digest 15  $\mu$ g of genomic DNA, and the digest was fractionated in 1% agarose gels. Southern blots were performed as described previously (36), and hybridization was carried out with a probe from the breakpoint region known from previous experiments not to contain repetitive DNA sequences.

#### RESULTS

For studies of globin mRNA expression during development, F<sub>1</sub> and F<sub>2</sub> progeny of the founders were utilized and RBCs for mRNA analysis from multiple transgenic animals from each line were obtained at desired stages of development. For a given developmental stage in a transgenic line, RNA samples from multiple fetuses and adult animals were analyzed to minimize experimental error. Only adult animals 5 weeks old or older (range, 5 weeks to 2 years) were utilized in these studies to prevent any bias resulting from analyzing adult animals shortly after birth for  $\gamma$ -globin gene expression.

**HPFH-2 3' breakpoint DNA sequences modify the developmental regulation of the  $\gamma$ -globin genes in transgenic mice in the absence of the LCR.** Two transgenic mouse lines carrying the G $\gamma$  transgene (lines 310 and 829) and two lines with the G $\gamma$ HPFH-2 transgene (lines 63 and 332) were established.

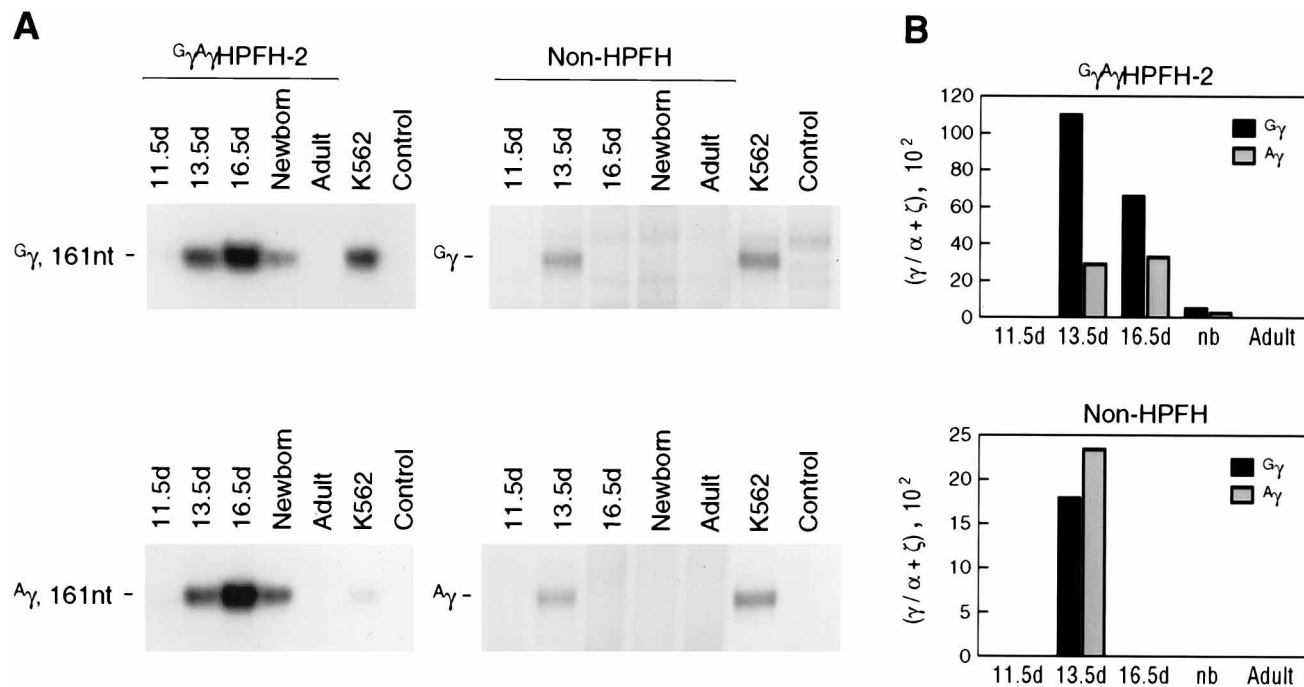


FIG. 2. (A) Human  $\gamma$ -globin mRNA expression in RBCs of transgenic mice carrying the  $G\gamma^A\gamma$ HPFH-2 transgene (line 332) and control (non-HPFH)  $G\gamma^A\gamma$  transgene (line 310) without the LCR. Ten micrograms of total RNA isolated from mouse RBCs at each indicated developmental stage was hybridized to  $G\gamma$ - or  $A\gamma$ -globin riboprobes in RNase protection assays as described by Starck et al. (40). The expected position of the specific 161-nucleotide (nt)  $G\gamma$  or  $A\gamma$  protected RNA fragment is indicated. Total RNA from human K562 cells and nontransgenic mouse blood RNA (Control) served as positive and negative controls, respectively. (B) Quantitative representation of the ratios of human  $G\gamma$  or  $A\gamma$  mRNA to murine  $\alpha$  plus  $\zeta$  mRNAs at each developmental stage. Results of HPHF-2 line 332 are shown on the top, and results of non-HPFH line 310 are shown on the bottom. Expression of the murine  $\alpha$ - plus  $\zeta$ -globin mRNAs was quantitated in separate experiments using probes with different specific activities and 100 ng of total RNA in each hybridization (data not shown). Therefore, vertical-axis values should be regarded not as absolute values but as arbitrary units of the ratios of human to murine mRNAs. nb, newborn.

Total RNA isolated from blood cells of mouse embryos at day 11.5, from 13.5- and 16.5-day fetuses, and from newborn pups and adult animals was analyzed for the presence of human mRNAs. Autoradiographs of representative RNase protection assays of lines 332 (HPFH-2) and 310 (non-HPFH) are shown in Fig. 2A. Expression of  $G\gamma$ - and  $A\gamma$ -globin mRNAs was detected at day 13.5 in both HPHF-2 (line 332) and non-HPFH (line 310) lines. After day 13.5, however, the pattern of  $\gamma$ -globin mRNA expression during development differed significantly in HPHF-2 and non-HPFH mice. At day 16.5, levels of the two  $\gamma$  mRNAs decreased markedly in mice carrying the  $G\gamma^A\gamma$  transgene (Fig. 2A). In contrast,  $\gamma$  mRNA expression is detected in 16.5-day fetuses as well as newborn mice carrying the  $G\gamma^A\gamma$ HPFH-2 transgene (Fig. 2A). Quantitative analysis of these results is illustrated in Fig. 2B. The ratios of  $G\gamma$ - and  $A\gamma$ -globin mRNAs relative to murine  $\zeta$  plus  $\alpha$  mRNAs for each stage of development were determined. Because the level of expression of the murine globin genes is markedly greater than that of the human transgenes in the absence of the LCR, human and murine globin mRNAs were analyzed on different gels with different amounts of total cellular RNA as described by Starck et al. (40). Therefore, the quantitative data illustrated in Fig. 2B represent not absolute values but relative arbitrary units of the ratio between human and murine globin gene expression at each stage of development. Similar results were obtained with lines 63 and 829 (data not shown). Thus, both  $G\gamma$ - and  $A\gamma$ -globin mRNAs were detected in RBCs of 16.5-day fetuses as well as newborn pups in the two HPHF-2 mouse lines, in contrast to the non-HPFH control mice carrying the  $G\gamma^A\gamma$  transgene, where levels of  $\gamma$ -globin mRNAs decreased markedly by fetal day 16.5 and were not detectable in

RBCs of newborn pups. Although no  $\gamma$ -globin gene expression was observed in RBCs of adult animals carrying the  $G\gamma^A\gamma$ HPFH-2 transgene without the LCR, these preliminary results indicated that DNA sequences located 3' to the breakpoint of the HPHF-2 deletion could modify the developmental expression of the  $\gamma$ -globin genes in transgenic mice and may be involved in the generation of the HPHF phenotype.

**The  $\gamma$ -globin genes are persistently expressed in erythroid cells of adult transgenic mice carrying the LCR $G\gamma^A\gamma$ HPFH-2 transgene.** Because our initial experiments with transgenic mice carrying the  $G\gamma^A\gamma$ HPFH-2 transgene without LCR showed  $\gamma$ -globin gene expression later during development than control  $\gamma$  transgenes but not in adult erythroid cells, we next wished to examine the effect of the LCR on  $\gamma$ -globin gene expression and asked whether the LCR is required to reproduce the full HPHF phenotype in transgenic mice. We analyzed, by quantitative RNase protection assays, six transgenic lines carrying the 30-kb LCR $G\gamma^A\gamma$ HPFH-2 transgene for expression of human  $\gamma$ - and endogenous mouse  $\alpha$ - and  $\zeta$ -globin mRNAs during mouse development. An autoradiograph of a representative RNase protection assay of line 3012 is shown in Fig. 3. Following PhosphorImager quantitation, human  $\gamma$ -globin mRNA levels were expressed as a percentage of mouse  $\alpha$ - and  $\zeta$ -globin mRNA levels and then corrected for the copy number of the transgenes as determined by Southern blotting. The results of  $\gamma$ -globin gene expression in the six transgenic mouse lines carrying the LCR $G\gamma^A\gamma$ HPFH-2 transgene are shown in Table 1. During the fetal stage of development, the highest level of human  $\gamma$ -globin gene mRNA per transgene copy ranged from  $7.3\% \pm 1.2\%$  to  $29\% \pm 6\%$  of the total mouse  $\alpha$ - plus  $\zeta$ -globin mRNA level. In erythroid cells of all

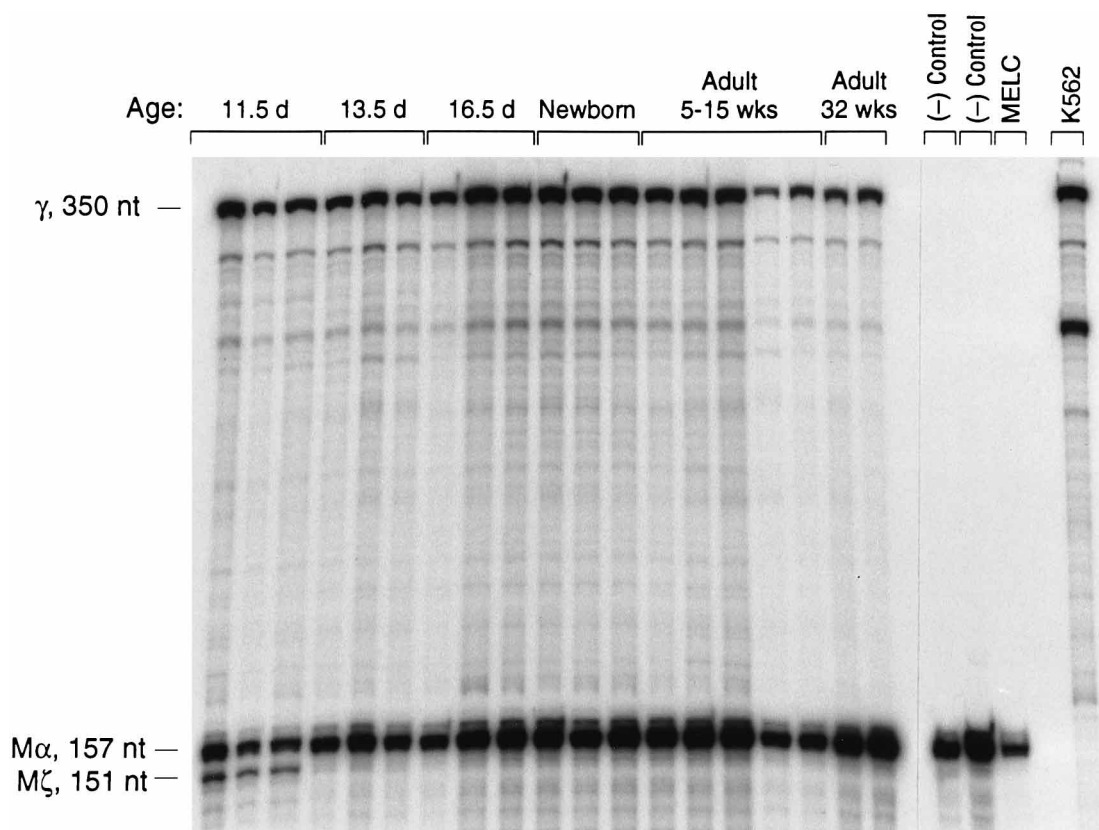


FIG. 3. Human  $\gamma$ -globin mRNA expression in HPFH-2 line 3012 carrying the  $LCR^{G\gamma^A\gamma}$ HPFH-2 transgene. Total RNA, isolated from blood of multiple individual animals at each developmental stage as indicated at the top of the figure, was hybridized simultaneously to human  $\gamma$ -, mouse  $\alpha$ -, and mouse  $\zeta$ -globin riboprobes (see Materials and Methods for the specific riboprobes and RNA quantities used for hybridization). The expected positions of the specific 350-nucleotide (nt) (human  $\gamma$ ), 157-nt (mouse  $\alpha$ ), and 151-nt (mouse  $\zeta$ ) protected RNA fragments are indicated. The positive control samples were total RNA isolated from mouse erythroleukemia cells (MELC) and human K562 cells. The negative (-) control samples were RNA extracted from blood of 13.5-day fetal and adult nontransgenic mice.

adult mice carrying the  $LCR^{G\gamma^A\gamma}$ HPFH-2 transgene, persistent expression of  $\gamma$ -globin mRNA was present, ranging from  $3.4\% \pm 0.2\%$  to  $8\% \pm 3.5\%$  per transgene copy of endogenous mouse  $\alpha$ -globin mRNA. Thus, as illustrated in Fig. 4, in adult animals, the expression of the  $\gamma$ -globin genes declined to only about one-third the level of maximal expression during fetal development, with an average percentage of adult  $\gamma$  mRNA level relative to the highest fetal  $\gamma$  mRNA level of  $34\% \pm 11\%$  (range, 21 to 48%). The decline in the level of  $\gamma$ -globin mRNA during development occurred predominantly prior to birth, with only a relatively minor decline after the newborn period, i.e., during the first 5 weeks following birth. The  $\gamma$ -globin mRNA levels remained quite stable up to 2 years of age in all adult animals tested (data not shown). Treatment with phenylhydrazine did not have an effect on  $\gamma$ -globin mRNA levels (data not shown). In all transgenic lines except line 3035,  $\gamma$ -globin gene expression was maximal at fetal day 11.5. Line 3035, which has 20 copies of the transgene, showed the highest  $\gamma$ -globin mRNA levels in 16.5-day fetuses in repeated experiments (Table 1). The results of  $\gamma$ -globin gene expression in six different transgenic lines carrying the  $LCR^{G\gamma^A\gamma}$ HPFH-2 transgene lead us to conclude that HPFH-2 breakpoint DNA sequences can modify the developmental pattern of  $\gamma$ -globin gene expression, resulting in significant levels of  $\gamma$ -globin mRNA in adult RBCs. Since our experiments using the 25-kb  $LCR^{G\gamma^A\gamma}$ HPFH transgene without the LCR did not reveal persistent  $\gamma$ -globin gene expression in adult erythroid cells, we

further conclude that the LCR is required for generation of the HPFH phenotype in transgenic mice.

Distribution of  $\gamma$ -globin chains in RBCs was examined by staining of peripheral blood with monoclonal anti- $\gamma$ -chain antibodies. Figure 5 shows representative slides. The distribution of  $\gamma$ -globin chains in adult RBCs was not pancellular, but overall a high percentage of the cells, ranging from 38 to 86%, stained positive.

**The  $\gamma$ -globin genes are silenced in RBCs of adult transgenic mice carrying the control  $LCR^{G\gamma^A\gamma}$  and  $LCR^{G\gamma^A\gamma\psi\beta}$  transgenes.** To serve as controls for the experiments carried out with the HPFH transgenic mice, one transgenic line with the 18-kb  $LCR^{G\gamma^A\gamma}$  transgene and one line with the 32-kb  $LCR^{G\gamma^A\gamma\psi\beta}$  transgene were generated. We asked whether  $\gamma$ -globin gene silencing could occur in the absence of a linked  $\beta$ -globin gene or HPFH-2 breakpoint DNA sequences in transgenic mice. Studies of gene expression during development by RNase protection assays were carried out as described above. A representative autoradiograph of an RNase protection gel for the transgenic line carrying the  $LCR^{G\gamma^A\gamma\psi\beta}$  transgene is shown in Fig. 6. Table 2 summarizes the results of  $\gamma$ -globin gene expression during development for each control line. In the line carrying the control  $LCR^{G\gamma^A\gamma}$  transgene, the highest  $\gamma$ -globin mRNA level during the fetal stage of development was  $31\% \pm 2.7\%$  (per transgene copy) of mouse  $\alpha$ - plus  $\zeta$ -globin mRNA, and the level declined significantly to  $0.4\% \pm 0.2\%$  in adult animals. Similarly, in the line carrying the  $LCR^{G\gamma^A\gamma\psi\beta}$  trans-

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