Histone H4 Hyperacetylation Precludes Histone H4 Lysine 20 Trimethylation*

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Posttranslational modification of histones is a common means of regulating chromatin structure and thus diverse nuclear processes. Using a hydrophilic interaction liquid chromatographic separation method in combination with mass spectrometric analysis, the present study investigated the alterations in histone H4 methylation/acetylation status and the interplay between H4 methylation and acetylation during in vitro differentiation of mouse erythroleukemia cells and how these modifications affect the chromatin structure. Independently of the type of inducer used (dimethyl sulfoxide, hexamethylenebisacetamide, butyrate, and trichostatin A), we observed a strong increase in non- and monoacetylated H4 lysine 20 (H4-Lys20) trimethylation. An increase in H4-Lys20 trimethylation, however, to a clearly lesser extent, was also found when cells accumulated in the stationary phase. Since we show that trimethylated H4-Lys20 is localized to heterochromatin, the increase in H4-Lys20 trimethylation observed indicates an accumulation of chromatin-dense and transcriptionally silent regions during differentiation and during the accumulation of control cells in the stationary phase, respectively. When using the deacetylase inhibitors butyrate or trichostatin A, we found that H4 hyperacetylation prevents H4-Lys20 trimethylation, but not mono- or di-acetylation, and that the nonacetylated unmethylated H4-Lys20 is therefore the most suitable substrate for H4-Lys20 trimethylase. Summarizing, histone H4-Lys20 hypotrimethylation correlates with H4 hyperacetylation and H4-Lys20 hypertrimethylation correlates with H4 hypoacetylation. The results provide a model for how transcriptionally active euchromatin might be converted to the compacted, transcriptionally silent heterochromatin.

In eukaryotes, histone proteins associate with DNA to form nucleosomes that are folded into higher order chromatin structures. Differences in higher order chromatin structures, which are important prerequisites for numerous biological processes including cellular proliferation, differentiation, development, gene expression, genome stability, and cancer, are thought to be realized by a variety of posttranslational modifications of histone N termini, particularly of histones H3 and H4. Besides acetylation, histones are subjected to phosphorylation, methylation, ubiquitination, ADP-ribosylation, and deamidation (1, 2). Distinct combinations of covalent histone modifications including lysine acetylation, lysine and arginine methylation, and serine phosphorylation form the basis of the histone code hypothesis (3–5). This hypothesis proposes that a pre-existing modification affects subsequent modifications on histone tails and that these modifications generate unique surfaces for the binding of various proteins or protein complexes responsible for higher order chromatin organization and gene activation and inactivation. Some of the histone-modifying enzymes (e.g. lysine methyltransferases) are, when deregulated, considered to be involved in carcinogenesis (6).

Histone H4 is typically acetylated at lysines 5, 8, 12, and 16, methylated at arginine 3 and lysine 20, and phosphorylated at serine 1 (3, 7, 8). Unlike the dynamic process of histone acetylation and phosphorylation, histone methylation is regarded as a relatively static long-term signal with a low turnover of the methyl group. Whereas arginine can be either mono- or dimethylated (the latter in symmetric or asymmetric form), lysine methylation can occur as a mono-, di-, or trimethylated derivative. In contrast to histone H3 methylation, H4-Lys20 was long considered to be maximally dimethylated in mammals (9). Most recently, however, Sarg et al. (10) conducted a mass spectrometric analysis with a newly developed hydrophilic interaction chromatographic method enabling the simultaneous separation of methylated and acetylated forms and, for the first time, found in vivo evidence that H4-Lys20 is also trimethylated in mammalian tissue. Moreover, in rat liver and kidney the proportion of trimethylated histone H4 increases during aging. In Raji and K562 cells the trimethylated form was also detected, primarily when the cells were accumulated in the stationary phase (10). Most recently, two novel SET domain histone methyltransferases, Suv4-20h1 and Suv4-20h2, acting as nucleosomal H4-Lys20 trimethylating enzymes, have been identified (11). It was also shown that histone H3-Lys9 trimethylation is required for the induction of H4-Lys20 trimethylation (H4-tri-mLys20) and that trimethylation of histone H3-Lys9 and histone H4-Lys20 functions as a repressive mark in gene-silencing mechanisms (11, 12).

Studies investigating possible links between histone methylation and acetylation have revealed that methylation of Arg2 of H4 by the histone H4-specific protein methyltransferase PRMT 1 facilitates acetylation by p300 on histone H4-Lys8 and -Lys12 and that this methylation at position 3 plays an important role in transcriptional regulation (7, 8). Acetylation on any of the four H4 lysines on the N-terminal tail (Lys5, Lys8, Lys12, Lys16), however, inhibits the methylation of Arg2 by PRMT 1 (7, 8). These results demonstrating the interplay between histone

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H4 methylation and acetylation support the histone code hypothesis. Recent findings suggest that histone H4-Lys16 acetylation inhibits H4-Lys16 acetylation and vice versa, leading to the hypothesis that methylation of H4-Lys20 maintains silent chromatin, in part by precluding neighboring acetylation on the H4 tail (13, 14). These studies, however, did not discriminate between the individual methylation states (3–5).

Of particular interest are recent findings regarding the role of various states of histone H3/H4 methylation in the regulation of transcription. In contrast to the dimethylated state of histone H3-Lys4, which occurs at both inactive and active euchromatic genes, H3-Lys9 trimethylation is present exclusively at active genes (15), whereas H3-Lys9 trimethylation and also H4-Lys20 trimethylation are present in repressive chromatin domains (11, 15). It is obvious, therefore, that not only methylation in itself but the precise methylation state and the position of the lysine determine gene activity/repression.

In the present study we aimed to investigate the changes in Lys20 methylation states of non-, mono-, di-, tri-, and tetraacetylated H4 histones as well as the interplay between the various H4-Lys20 methylation states and acetylation during in vitro differentiation of mouse erythroleukemia cells induced by Me2SO/HMBA and the deacetylase inhibitors sodium butyrate/TSA, respectively. For the analysis we used a hydrophilic interaction liquid chromatographic method recently developed in our laboratory (2, 16–19) in combination with a mass spectrometric analysis enabling the simultaneous separation of non-, mono-, di-, and trimethylated Lys20 of non-, mono-, di-, and tetraacetylated H4 histones. In all cases and regardless of the nature of the inducer we found a strong increase in the trimethylated Lys20 of non- and monoacetylated histone H4 (ac0 H4-tri-mLys20 and ac1 H4-tri-mLys20, respectively) during differentiation. By using an antibody that specifically recognizes trimethylated Lys20 of histone H4, it was possible to demonstrate that H4-tri-mLys20 localizes to heterochromatin. The observed increase in non- and monoacetylated H4-tri-mLys20 thus results in an increase in heterochromatic regions, thereby indicating chromatin compaction and repression of gene activity in the course of erythroid differentiation. Treatment of mouse erythroleukemia cells with sodium butyrate or TSA resulted in H4 histone hyperacetylation. With regard to the individual acetylated forms in detail, we found diacetylated H4-Lys20 to be trimethylated in traces, whereas tri- and tetraacetylated H4-Lys20 were not trimethylated at all. We therefore suggest that in contrast to non- and monoacetylated H4 histones, which correlate with H4-tri-mLys20, histone H4 hyperacetylation precludes Lys20 trimethylation, thus providing further support for the histone code hypothesis. Independently of the degree of acetylation, however, we found histone H4-di-mLys20 and, to a far lesser extent, histone H4-mono-mLys20

**EXPERIMENTAL PROCEDURES**

**Materials**—Sodium perchlorate (NaClO4), triethylamine (TEA), acetanilide, and trifluoroacetic acid were purchased from Fluka (Buchs, Switzerland). All other chemicals were purchased from Merck (Darmstadt, Germany) if not otherwise indicated.

**Cell Culture**—Mouse erythroleukemia (MEL) cells (line F4N) were grown in Dulbecco's minimum Eagle's medium (Biochrom, Berlin, Germany) containing 2% nonessential amino acids, 1% penicillin-streptomycin, and 10% fetal calf serum. Cells were cultured at initial cell densities of 5.104–7.104/ml at 37 °C and 5% CO2. Differentiation was induced by the addition of 2% MeSO4 (for 144 h), 5 mM HMBA (144 h), 4 ng/ml TSA (144 h), or 1.75 mM sodium butyrate (72 h). The untreated control cells were harvested after 72 h in the log phase or after 144 h in stationary phase. The percentage of benzidine-positive cells was determined as described by Orkin et al. (20).

**Preparations of Histones**—The histones were extracted from MEL cells with sulfuric acid (0.2 M) according to the procedure of Helliger et al. (21).

**High Performance Liquid Chromatography**—The equipment used consisted of a 127 Solvent Module and a Model 166 UV-visible region Detector (Bockman Instruments). The effluent was monitored at 210 nm, and the peaks were recorded using Beckman System Gold software.

**Reversed-phase HPLC**—The separation of core histones was performed on a Nucleosil C4 column (250 × 8 mm inner diameter, 5-μm particle pore size, 30-nm pore size, end-capped; Seibersdorf). Samples (20 μl) were injected using an Agilent 1100 sample manager. The histone sample was chromatographed as described previously (10, 22).

**Hydrophilic Interaction Liquid Chromatography**—The histone fraction H4 (~100 μg) isolated by RP-HPLC was further separated on a SynChropak CM300 column (250 × 4.6-mm inner diameter, 6.5-μm particle size, 30-nm pore size; Agilent Technologies, Vienna, Austria) at 30 °C and at a constant flow of 1.0 ml/min using a multi-step gradient (solvent A: 100% (v/v) solvent A (0.067% 2-mercaptoethanol, 0.015% TEA/H3PO4, pH = 3.0; solvent B: 65% acetonitrile, 0.015% TEA/H3PO4, pH = 3.0, and 0.68 mM NaClO4). The concentration of solvent B was increased from 0 to 10% in 2 min and from 10 to 40% in 30 min and was then maintained at 40% for 10 min. The isolated protein fractions were desalted using RP-HPLC. Histone fractions obtained in this way were collected and, after the addition of 20 μl of 2-mercaptoethanol (0.2 M), were lyophilized and stored at −20 °C.

**Endoprotease Glu-C Digestion**—H4 histone (~10 μg) obtained by HILIC fractionation was digested with Staphylococcus aureus V8 protease (Roche Applied Science; 1:20 w/w) in 50 μl of 25 mM NH4HCO3 buffer (pH 4.0) for 1 h at room temperature. The digest was subjected to RP-HPLC as described previously (10).

**Endoprotease Lys-C Digestion**—N-terminal peptides obtained by Glu-C digestion were further digested with endoprotease Lys-C (Roche Applied Science; 1:5 w/w) in 15 μl of 25 mM Tris-HCl buffer (pH 8.2) for 2 h at 37 °C. The digest was subjected to RP-HPLC-ESI-MS.

**Mass Spectrometric Analysis**—Histone H4 peptide fractions obtained by endoprotease Lys-C cleavage were injected onto a Nano-PepMap C18 column (150 × 1-mm inner diameter, 3-μm particle size; ICT, Vienna, Austria). The column eluate was coupled directly to a Finnigan LCQ ion trap mass spectrometer equipped with an electrospray ion source (RP-HPLC-ESI-MS). Samples of ~1 μg were chromatographed within 55 min at a constant flow of 35 μl/min with a two-step acetonitrile gradient starting at solvent A − solvent B (90:10) (solvent A: water containing 0.1% trifluoroacetic acid; solvent B: 85% acetonitrile and 0.983% trifluoroacetic acid). The concentration of solvent B was increased linearly from 10 to 40% over a 45-min period and from 40 to 100% over 20 min.

**Antibodies**—The antibodies used were: rabbit polyclonal to H4-tri-mLys20 (Abcam), Western blotting 1:1000, immunofluorescence 1:200; rat monoclonal to HPIβ (Abcam), immunofluorescence 1:50; mouse monoclonal to RNA polymerase II (Abcam), immunofluorescence 1:500. For Western blotting, goat anti-rabbit IgG peroxidase conjugate (Sigma) 1:5000 was used as secondary antibody. The secondary antibodies for immunofluorescence were: goat anti-rabbit IgG fluorescein isothiocyanate conjugate (Sigma), 1:160; goat anti-rat IgG Cy3 conjugate (Jackson ImmunoResearch), 1:160; goat anti-mouse IgG Cy3 conjugate (Jackson ImmunoResearch), 1:160.

**Western Blotting**—Histones were resolved in SDS–loading buffer, fractionated by SDS–PAGE, and transferred to a nitrocellulose membrane (Hybond ECL, Amersham Biosciences). The membrane was probed with antibodies using standard techniques and detected by enhanced chemiluminescence (ECL, Amersham Biosciences).

**Immunofluorescence**—MEL cells were grown in suspension culture, collected by centrifugation, and washed with phosphate-buffered saline. 106 cells were cytospun onto slides. Cells were permeabilized using Triton X-100 (0.1% for 2 min and 0.01% for 10 min) in KCM buffer (120 mM KCl, 20 mM NaCl, 0.5 mM EDTA, 10 mM Tris/HCl, pH 7.5) and incubated sequentially with the primary and secondary antibodies. Cells were fixed for 5 min with 4% paraformaldehyde in KCM, stained with DAPI for 1 min, and mounted in mounting medium (Dako). Staining was visualized using an ×100 objective on a Zeiss Axioskop 2 microscope and a SPOT camera (Diagnostic Instruments). Images were captured using MetaVue software (Universal Imaging Corp.) and analyzed with Corel Photo-Paint 10.
RESULTS

HILIC Separation of Modified Forms of Histone H4 of Uninduced MEL Cells and Cells Cultured in the Presence of Me2SO, HMBA, Sodium Butyrate, and TSA—Untreated MEL cells and cells treated with Me2SO, HMBA, sodium butyrate, and TSA were grown in culture. After 72 and 144 h and before harvesting the cells, the number of benzidine-positive cells indicating hemoglobin accumulation was determined. In the case of butyrate-treated cells, however, the study was limited to the first 72 h, because butyrate is metabolized by the cells, and after a certain time its concentration is reduced to a level below that required to stimulate differentiation (23). Whereas throughout the entire period study untreated cultures were found to contain a very low level (less than 2%) of spontaneously differentiated cells, cultures containing Me2SO, HMBA, or butyrate produced 80–95% benzidine-positive cells and those containing TSA nearly 70%.

The sulfuric acid-extracted core histones were fractionated using an RP-HPLC procedure described previously (10, 22), and the histone H4 fractions obtained were subjected to HILIC, a chromatographic technique developed in our laboratory for separating modified core and H1 histones (2, 16–19). A number of major and minor peaks could be received, which were identified by ESI-MS as described previously (10). Fig. 1 shows the pattern of histone H4 acetylation and methylation of untreated (control) MEL cells at 72 h (Fig. 1A) and 144 h (Fig. 1B) after seeding, respectively, and of cells grown for 72 h in the presence of butyrate (Fig. 1C) and 144 h in the presence of Me2SO (Fig. D). The histone H4 acetylation/methylation pattern of MEL cells grown for 144 h in the presence of TSA and HMBA resembled that obtained by treating with butyrate and Me2SO, respectively (data not shown). In control cells and cells treated with Me2SO and HMBA, histone H4 was hypoacetylated, containing primarily non acetylated (ac0) and monoacetylated (ac1) forms with Lys16 as the most frequently acetylated lysine residue. In response to Me2SO treatment, a clear increase in trimethylated unacetylated and trimethylated monoacetylated H4 species was observed (Fig. 1, D versus B). Treatment of cells with butyrate and TSA, known HDAC inhibitors, caused hyperacetylation of H4, in which up to four acetyl groups are bound to the N-terminal tail. The results revealed that H4-Lys20 is methylated in all of the acetylated isoforms, as well as in the nonacetylated isoform, apart from a nonmethylated H4 present in varying amounts. In general, the dimethylated form was found to be the main methylation product followed, in smaller amounts, by the mono- and trimethylated forms. To facilitate an unambiguous assignment, the double peak of triacetylated H4 histones (designated in Fig. 1C with an asterisk) had to be investigated in more detail.

Characterization of Hyperacetylated H4 Fractions from Butyrate-treated Cells Obtained by HILIC—To precisely deter-
H4 Hyperacetylation Precludes H4-Lys\(^{20}\) Trimethylation

![Image](image-url)

**FIG. 2.** Peptide patterns obtained after endoproteinase Glu-C and Lys-C digestion of HILIC-Peak ac3 (marked with an asterisk). The peak marked with an asterisk, which was obtained from butyrate-treated MEL cells by HILIC (Fig. 1C), was digested with endoproteinase Glu-C. The resulting peptide fractions were separated using RP-HPLC (data not shown). The blocked N terminus (residues 1–52, 1–53) obtained was further digested by Lys-C. The resulting peptides and their attached modifications were identified using ESI-MS. Mouse histone H4 sequence data were taken from the Swiss Protein Database (accession number P02304).

mine the modification status of hyperacetylated histone H4 shown in Fig. 1C, the HILIC fractions were isolated, desalted by RP-HPLC, and treated with endoproteinase Glu-C as described previously (10). The fragmentation yielded an N-terminal peptide (residues 1–52 and 1–53) that was further cleaved with endoproteinase Lys-C. Endoproteinase Lys-C is a serine protease, and its activity is inhibited at lysine residues that have been modified by an acetyl or methyl group (10). The digests were analyzed by RP-HPLC-ESI-MS using a 150 × 1.0-mm inner diameter microbore column. The diacylated (ac2) form was found to be acetylated at Lys\(^{16}\) and Lys\(^{12}\), and as can be seen from the separation pattern in Fig. 1C, it was mainly dimethylated followed by the monomethylated derivative and, present in trace amounts only, the trimethylated one. In the case of the triacylated H4 protein (marked with an asterisk), which is separated into two HILIC subfractions, several peptide fragments were detected (Fig. 2). ESI-MS analysis revealed a triacylated H4-di-meLys\(^{20}\) form with acetyl groups at either Lys\(^{5}\)-Lys\(^{12}\)-Lys\(^{16}\) or Lys\(^{5}\)-Lys\(^{12}\)-Lys\(^{16}\) and, in very small amounts, at Lys\(^{5}\)-Lys\(^{8}\)-Lys\(^{16}\). This finding that the various lysines are partially occupied in the triacylated form agrees well with observations made by other investigators (24, 25). Interestingly, a peptide corresponding to the tetraacylated (Lys\(^{5}\)-Lys\(^{8}\)-Lys\(^{12}\)-Lys\(^{16}\)) nonmethylated form, which could not be separated by the HILIC system, was found in the ac3 double peak. A triacylated trimethylated H4 histone, however, was undetectable. Concerning the tetraacylated H4 protein, two HILIC subfractions containing one or two methyl groups were identified. Similar to triacylated H4, a tetraacylated trimethylated H4 was also not seen. At any rate, however, the H4-di-meLys\(^{20}\) form predominates in the hyperacetylated histone H4 fractions.

The Levels of H4-tri-meLys\(^{20}\) Are Raised in Hypoacetylated H4 Fractions during MEL Cell Differentiation—We recently showed for the first time that histone H4 from mammalian tissue is not only mono- and dimethylated but also trimethylated at Lys\(^{20}\) and that the trimethylated form accumulates with age and in cells in the stationary phase (10). Novel studies on methylated histones have revealed that H4-tri-meLys\(^{20}\) is a repressive mark in gene silencing and is important for heterochromatin assembly (11, 12). We were thus interested in investigating the occurrence of trimethylated H4 in differentiated cells. To obtain quantitative data, we conducted a HILIC analysis of histone H4 from MEL cells treated with various inducers and quantified the results. As depicted in Fig. 3A, an increase of about 70% in trimethylated H4 (ac0 + ac1) forms appears in non-growing cells. Upon induction a remarkable further increase to the 4–5-fold level occurs in differentiated cells. In addition, using an anti-H4-tri-meLys\(^{20}\) antibody we investigated the level of H4-tri-meLys\(^{20}\) in cells treated with Me\(_{6}\)SO for 96 h (Fig. 3B, lane 4) and in untreated control cells (Fig. 3B, lane 3). In Me\(_{6}\)SO-induced cells we found H4-tri-meLys-increase similar to that observed in HILIC (Fig. 3A). Fig. 3C shows the relative amount of H4-tri-meLys\(^{20}\) from the nonacylated fraction in comparison with the monoacylated one. Besides the increase in H4-tri-meLys\(^{20}\) in both the ac0 and ac1 fraction of treated and untreated cells, it is obvious that the proportion of trimethylation is always about 2-fold higher in nonacylated than in monomethylated H4. Nishioka et al. (13) found that in Drosophila, methylation of H4-Lys\(^{20}\) and acetylation of Lys\(^{16}\) are competitive. We favor the view that not methylation in itself but the degree of methylation plays a role in that interplay and that acetylated H4-Lys\(^{16}\) does not influence mono- or dimethylation but reduces trimethylation of H4-Lys\(^{20}\). It is also interesting to note that in non- and monomethylated H4 with increasing trimethylation we saw a decreasing amount of H4-non-meLys\(^{20}\) (designated as 0 in Fig. 1). This observation is in line with the in vitro finding that Suv4-20h is a Lys\(^{20}\) trimethylase preferring nonmethylated H4 as substrate (11).

H4-tri-meLys\(^{20}\) Localizes to Heterochromatin—To investigate the localization and distribution of H4-tri-meLys\(^{20}\) in chromatin of untreated control and differentiated MEL cells we used indirect immunofluorescence analysis (Fig. 4). Generally, during in vitro differentiation of MEL cells, their nuclei were reduced in volume and the chromatin appeared to be highly compacted (Fig. 4, lower row in both A and B, DAPI stain). Important markers of condensed DNA, which are key components of constitutive heterochromatin, represent the heterochromatin proteins HP1\(_{a}\) and HP1\(_{b}\) (26). We found that H4-tri-meLys\(^{20}\) was enriched mainly within DAPI-dense regions, which almost completely overlap with HP1\(_{a}\)-stained chromatin regions both in control and in Me\(_{6}\)SO-treated cells (Fig. 4A). To verify this finding and to exclude any association between H4-tri-meLys\(^{20}\) and transcriptionally engaged chromatin, dual staining experiments were performed using the H4-tri-meLys\(^{20}\)-specific antibody and an antibody against RNA polymerase II (27). We found that the majority of H4-tri-meLys\(^{20}\) is
Histone H4 Hyperacetylation Precludes H4-Lys²⁰ Trimethylation

In this study we aimed to quantify the changes in histone H4 methylation and acetylation levels in the course of in vitro differentiation of mouse erythroleukemia cells and to investigate their consequences regarding heterochromatin formation. Using a polar stationary phase in the presence of acetonitrile, unmodified H4 and its differently acetylated/methylated forms were separated from each other due to slight variations in the polarity of the proteins caused by the modification reactions. To emphasize the polar nature of intermolecular forces governing retention with such stationary phases, the term hydrophilic interaction chromatography, with the acronym HILIC, was suggested by Alpert (29). This HILIC-MS technique applied has a series of advantages over immunoassays now commonly used in this field. First, the whole histone H4 acetylation and methylation pattern is obtained not only simultaneously but also in extreme detail. In addition, the method requires only small sample amounts and gives results within 35 min.

When untreated cells were accumulated in the stationary phase, the proportion of non- and monoacetylated H4-tri-meLys²⁰ increased modestly, whereas in particular that of the diacetylated H4-tri-meLys²⁰ was found in each of them. Concerning the diacetylated H4 fraction, the smallest traces of H4-tri-meLys²⁰ might have been present in some samples treated with Me₂SO, HMBA, and TSA. Minimal but detectable amounts, however, were present in samples from butyrate-treated cells. The mono- and dimethylated form of Lys²⁰ was also seen in the hyperacetylated histone H4 fractions, but trimethylated Lys²⁰ was not detected in the tri- and tetraacetylated proteins. As mentioned before, Lys¹⁰ acetylation causes a decrease in Lys²⁰ trimethylation, whereas hyperacetylation with further acetyl groups at Lys¹², Lys⁸, and/or Lys⁶ even causes a complete repression of H4-Lys²⁰ trimethylation.

DISCUSSION

In this study we aimed to quantify the changes in histone H4 methylation and acetylation levels in the course of in vitro differentiation of mouse erythroleukemia cells and to investigate their consequences regarding heterochromatin formation. Using a polar stationary phase in the presence of acetonitrile, unmodified H4 and its differently acetylated/methylated forms were separated from each other due to slight variations in the polarity of the proteins caused by the modification reactions. To emphasize the polar nature of intermolecular forces governing retention with such stationary phases, the term hydrophilic interaction chromatography, with the acronym HILIC, was suggested by Alpert (29). This HILIC-MS technique applied has a series of advantages over immunoassays now commonly used in this field. First, the whole histone H4 acetylation and methylation pattern is obtained not only simultaneously but also in extreme detail. In addition, the method requires only small sample amounts and gives results within 35 min.

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FIG. 3. Increase of H4-tri-meLys²⁰ of hypoacetylated (ac⁰ + ac¹) forms during MEL cell differentiation. A, H4 was obtained from untreated (control) MEL cells 72 and 144 h after seeding and from cells grown for 72 h in the presence of butyrate and for 144 h in the presence of dimethyl sulfoxide (DMSO), HMBA, and TSA. The H4 fractions (~100–300 µg) isolated with RP-HPLC were analyzed under the same HILIC conditions described in the legend for Fig. 2. The amount of H4-tri-meLys²⁰ was quantified using Beckman System Gold software. The relative increase in H4 trimethylation of nongrowing cells and induced cells was compared with the 72-h control (0%). B, MEL cells were cultured for 96 h in normal culture medium (lanes 1 and 3) or in culture medium containing 2% dimethyl sulfoxide (lanes 2 and 4). Histones from the two cell samples were separated by SDS-PAGE, blotted, and probed with H4-tri-meLys²⁰ polyclonal antibody. Lanes 1 and 2 show a Coomassie Blue-stained profile of the samples used in the immunoblot (lanes 3 and 4). C, same conditions as described in A. The relative amount of nonacetylated (ac⁰) H4-Lys²⁰ trimethylation was compared with the monomethylated (ac¹) trimethylated form. Results represent means ± S.D. for three to five independent experiments.
and, to a lesser degree, monoacetylated histone H4 and that H4 hyperacetylation and Lys\(^{20}\) trimethylation preclude one another completely. This result is consistent with earlier studies demonstrating that H4 methylation preferentially occurs on unacetylated histones (9). It should be pointed out, however, that the mutual inducement observed solely concerns the interplay between hyperacetylation and Lys\(^{20}\) trimethylation but not between hyperacetylation and Lys\(^{20}\) mono- or dimethylation. In fact, we found non-, mono-, di-, tri-, and tetraacetylated H4 histones that were mono- and dimethylated. This outcome also agrees well with the finding that H4-di-meLys\(^{20}\) is broadly

**FIG. 4.** Co-immunostaining of differentiated (treatment with dimethyl sulfoxide for 96 h) and untreated (Control) MEL cells using antibodies that detect HP1\(\beta\), RNA polymerase II, and H4-tri-meLys\(^{20}\). A, distribution of HP1\(\beta\) (red), which detects heterochromatic sites, and H4-tri-meLys\(^{20}\) (green) in mouse MEL nuclei. In the far right column, the HP1\(\beta\) (Cy3, red) and H4-tri-meLys\(^{20}\) (FITC, green) staining patterns were visualized simultaneously (Merge). DNA was counterstained with DAPI to highlight the foci of heterochromatin. B, distribution of RNA polymerase II (red), which detects transcriptionally competent regions, and H4-tri-meLys\(^{20}\) (green) in MEL nuclei. In the far right column, the polymerase II (Cy3, red) and H4-tri-meLys\(^{20}\) (FITC, green) staining patterns were visualized simultaneously (Merge). DNA was counterstained with DAPI to highlight the foci of heterochromatin. Scale bars, 5 \(\mu\)m.

**FIG. 5.** A proposed model for how epigenetic modifications might cause gene silencing. HDAC deacetylates lysines 5, 8, 12, and 16 in the N-terminal tail of histone H4, which can in turn be trimethylated at Lys\(^{20}\) by Suv4-20h. Trimethylated H4-Lys\(^{20}\) is recognized by HP1, resulting in the maintenance of gene silencing.

**TABLE I**

Acetylation/methylation status of histone H4

A summary of the occurrence of various acetylated and methylated forms in untreated (control) cells and cells treated with several inducers is shown. Results show the presence (+) or absence (−) of mono-, di-, or trimethylated H4-Lys\(^{20}\) in the various acetylated H4 proteins.

| H4          | Controls | Me\_SO/HMBA | Butyrate/TSA |
|-------------|----------|-------------|--------------|
| ac\(0\)     |          |             |              |
| mono-me     | +        | +           | +            |
| di-me       | +        | +           | +            |
| tri-me      | +        | +           | +            |
| ac\(1\)     |          |             |              |
| (Lys\(^{16}\)) mono-me | + | + | + |
| di-me       | +        | +           | +            |
| tri-me      | +        | +           | +            |
| ac\(2\)     |          |             |              |
| (Lys\(^{16},Lys^{12}\)) mono-me | + | + | + |
| di-me       | +        | +           | +            |
| tri-me      | −        | −           | (+)          |
| ac\(3\)     |          |             |              |
| (Lys\(^{16},Lys^{12},Lys^{8}\)) mono-me | − | − | + |
| (Lys\(^{16},Lys^{12},Lys^{8}\)) di-me | − | − | + |
| (Lys\(^{16},Lys^{8}\)) tri-me | − | − | − |
| ac\(4\)     |          |             |              |
| (Lys\(^{16},Lys^{12},Lys^{8}\)) mono-me | − | − | + |
| di-me       | −        | −           | +            |
| tri-me      | −        | −           | −            |
distributed in euchromatin (11).

The fact that certain combinations of acetyl and methyl modifications of lysines in histone tails may have antagonistic or cooperative biological effects is repeatedly described in the literature (7, 9, 13, 14, 37–40). Although acetylation at Lys\(^3\) in histone H3 prevents methylation of that amino acid (37), acetylation at any of the four lysines (Lys\(^3\), Lys\(^4\), Lys\(^12\), Lys\(^16\)) of histone H4 inhibits Arg\(^5\) methylation, whereby acetylation at Lys\(^3\) has the greatest effect (7). In this context it should be mentioned that our study detected no methylation of H4-Arg\(^5\). Nishioka et al. (13) described the identification and characterization of a mammalian histone methyltransferase, PR-Set7, which specifically methylates the Lys\(^20\) of histone H4. In \textit{in vitro} experiments these authors (13) demonstrated that methylation at H4-Lys\(^20\) inhibits the acetylation of H4-Lys\(^16\) and that, vice versa, acetylation of H4-Lys\(^16\) precludes methylation of the neighboring Lys\(^20\). The acetylation of Lys\(^12\), however, was not found to be inhibited by Lys\(^20\) methylation. In this context it should be noted that the investigations of Nishioka et al. (13) were performed using dimethylated H4-Lys\(^20\), whereas our study revealed the interactions between acetylation and mono-, di-, and trimethylation of H4-Lys\(^20\).

The mechanisms by which euchromatin is converted to heterochromatin are still unknown. Based on the results obtained in our study, we propose a model showing how epigenetic modifications might convert unmethylated “active” chromatin into trimethylated “silenced” chromatin (Fig. 5). The first step involves the deacetylation of hyperacetylated H4 by specific HDACs. Thereafter, the unmodified histone H4 is trimethylated at Lys\(^3\) by a Suv4-20h methylase. If there exists an Suv4-20h- HDAC multienzyme complex similar to the Su(VAR3)-3-HDAC1 protein complex present in \textit{Drosophila} histone H3 (31), it would also be imaginable that deacetylation and trimethylation occur in a concerted manner. In the subsequent last step, HP1 binds to the H4-tri-meLys\(^20\), which in turn causes chromatin compaction.

Histone modifications and DNA methylation are the major epigenetic mechanisms that can effect gene expression in mammals (38). Recent studies indicate a connection between histone H3 and DNA methylation (39–42). Tamaru et al. (41) suggested that H3-tri-meLys\(^8\) affects DNA methyltransferases, causing cytosine methylation on nearby DNA. The observations support the idea that histone H3 and DNA methylation cooperate in establishing long-term states of transcriptional regulation. Because H3-Lys\(^8\) and H4-Lys\(^20\) trimethylation have the same functional consequences, we also assume an interaction between H4-tri-meLys\(^20\) and DNA methyltransferases.

It is known that DNA hypermethylation and histone deacetylation are involved in tumor suppressor gene silencing in cancer (41, 43, 44). For this reason, inhibitors of histone deacetylases, such as TSA, have great potential for the treatment of malignant disease (45). Recently, it was shown that histone H3-Lys\(^8\) methylation/acetylation also has a crucial function in epigenetic regulation of suppressor genes: combined treatment with TSA and a DNA methyltransferase inhibitor resulted in an increase in H3-Lys\(^8\) acetylation and a simultaneous decrease in histone H3-Lys\(^8\) methylation, correlating with re-expression of silenced suppressor genes (43, 44). We surmise that there is a similar TSA effect on histone H4. As our study shows, TSA, similar to sodium butyrate, causes H4 hyperacetylation, which prevents Lys\(^3\) trimethylation and then heterochromatin formation. In this context it would be interesting to know whether a H4 trimethyltransferase inhibitor would intensify the efficacy of deacetylase inhibitors.