Human genomic \(Z\)-DNA segments probed by the \(Z\alpha\) domain of ADAR1

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ABSTRACT

Double-stranded DNA is a dynamic molecule that adopts different secondary structures. Experimental evidence indicates \(Z\)-DNA plays roles in DNA transactions such as transcription, chromatin remodeling and recombination. Furthermore, our computational analysis revealed that sequences with high \(Z\)-DNA forming potential at moderate levels of DNA supercoiling are enriched in human promoter regions. However, the actual distribution of \(Z\)-DNA segments in genomes of mammalian cells has been elusive due to the unstable nature of \(Z\)-DNA and lack of specific probes. Here we present a first human genome map of most stable \(Z\)-DNA segments obtained with A549 tumor cells. We used the \(Z\)-DNA binding domain, \(Z\alpha\), of the RNA editing enzyme ADAR1 as probe in conjunction with a novel chromatin affinity precipitation strategy. By applying stringent selection criteria, we identified 186 genomic \(Z\)-DNA hotspots. Interestingly, 46 hotspots were located in centromeres of 13 human chromosomes. There was a very strong correlation between these hotspots and high densities of single nucleotide polymorphism. Our study indicates that genetic instability and rapid evolution of human centromeres might, at least in part, be driven by \(Z\)-DNA segments. Contrary to \textit{in silico} predictions, however, we found that only two of the 186 hotspots were located in promoter regions.

INTRODUCTION

Left-handed \(Z\)-DNA is an alternative secondary structure to the right-handed \(B\)-conformer (1). It represents a higher energy state with a short half life, unless stabilized by factors such as negative \((-)\) DNA supercoiling or chemical DNA modification (2,3). \(Z\)-DNA occurs preferentially in stretches of alternating purine/pyrimidine residues, where the energetic barrier that accompanies a \(B\)-to-\(Z\) transition is smallest (2). Potential biological functions of \(Z\)-DNA have been investigated for over 30 years, since the first molecular structure was revealed by X-ray crystallography (4). Experimental evidence points to the existence of \(Z\)-DNA in living mammalian cells (5) and a functional role in processes such as gene regulation (6,7), nucleosome positioning (8,9), chromatin remodeling (10) and recombination (11,12).

Whole genome mapping of \(Z\)-DNA has been limited to \textit{in silico} predictions (13–15), which showed that high potential \(Z\)-DNA forming regions (\(Z\)DRs) are located preferentially in close proximity to transcriptional start sites (TSS). This finding together with the fact that translocating RNA polymerases can induce \((-)\) supercoiling in their wake \textit{in vivo}, raised expectations that segments of \(Z\)-DNA exist near TSS in a number of gene regulatory regions during the cell cycle (2,3). However, direct evidence is lacking that this is a widespread phenomenon and, if so, that these \(Z\)DRs have a function.

We addressed this topic here and developed first a \(Z\)-DNA prediction program to determine genomic \(Z\)DRs that undergo structural transitions at different levels of DNA supercoiling. We then used the \(Z\alpha\) domain of the human RNA editing enzyme ADAR1 (\(Z\alpha\)ADAR1) as a \(Z\)-DNA-specific probe to obtain direct evidence for the existence of \(Z\)DRs in \(Z\)-conformation within human cells. \(Z\alpha\)ADAR1 recognizes \(Z\)-DNA through two distinct features: the zigzag phosphate backbone and the \textit{syn} conformation of one purine residue in a \(Z\)-DNA segment. The protein binding site occupies only 6 bp of \(Z\)-DNA, yet associates with high affinity to the \(Z\)-conformer of many different nucleotide sequences (16,17). Using this versatile probe in conjunction with a dual crosslinking chromatin affinity precipitation (ChAP) strategy, we present here a first map of \(Z\)-DNA segments in the human genome and discuss biological implications.

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MATERIALS AND METHODS

Construction and purification of probes

Za (GI:2795789) was amplified by PCR from pET28a-Za77 (17). The product was further amplified by PCR to add sequences encoding tags FLAG and Strep II to the 3′-end. The final product was digested with NdeI and EcoRI, and ligated into pET17b as described above. Mutations were introduced at N173A and Y177A by assembly PCR, and cloned again into pET17b as described above. The purification of probes is described in detail in Supplementary Data.

In vitro binding and crosslinking of Za to Z-DNA

Genomic DNA fragments containing a d(GT)$_6$, insert were amplified from the promoter region of the mouse mast cell protease 6 gene (18) and inserted into pPGKss-puro vector using EcoRI, which resulted in plasmid pGT-pPGKss-Puro. Total 1.5 μg of supercoiled pGT-pPGKss-puro was mixed with 200 ng of a 360 bp d(GT)$_6$ containing PCR fragment amplified from pGT-pPGKss-puro with primers GT-U 5′-CCCTTCTGATGACCACAGGTCAC 3′ and GT-L 5′-TCCAGACTGCTTTGGGAAAG3′. The DNA mixture was diluted with 350 μl of HEPES binding buffer (50 mM HEPES pH 7.4, 150 mM NaCl, 1 mM EDTA, 1 mM DTT) and incubated with 1 μg of purified Z$_{ADAR1}$, Z$_{ADAR1mut}$ or BSA at room temperature for 15 min. Total 0.5% formaldehyde was added and further incubated for 5 min, followed by an incubation with 125 mM glycine for additional 5 min. Ethanol precipitation was used to purify DNA, which was subsequently cleaved with EcoRI and analyzed by EMSA.

In vitro Chromatin Affinity Precipitation (ChAP)

About 2 × 10$^6$ A549 cells were crosslinked with formaldehyde and treated with Triton X-100 as described (10). Cells were washed twice with cold PBS and a buffer containing 50 mM HEPES, pH 8.0, 150 mM NaCl, 1 mM EDTA. Thirty micrograms of purified Z$_{ADAR1}$, Z$_{ADAR1mut}$ or BSA were diluted into 5 ml of HEPES-binding buffer and added to the cells. After incubation at 4°C for 5 h, cells were washed four times with HEPES-binding buffer at 4°C. For the second crosslinking reaction, 0.5% formaldehyde was added in 10 ml HEPES-binding buffer and incubated for 5 min at room temperature. Crosslinking was terminated as described above, and cells were washed five times with cold PBS and harvested in 4 ml cold ChAP lysis buffer (50 mM HEPES, pH 8.0, 1 mM EDTA, 0.5 mM EGTA, 140 mM NaCl, 10% glycerol, 0.5% NP-40, 0.25% Triton X-100, 1 mM PMSF). Nuclei were collected at 1500 g for 5 min at 4°C and resuspended in nuclei wash buffer (10 mM Tris–HCl, pH 8.0, 1 mM EDTA, 0.5 mM EGTA, 200 mM NaCl, 1 mM PMSF). Nuclei were harvested again and resuspended in 200 μl of SDS lysis buffer (50 mM Tris–Cl, pH 8.1, 10 mM EDTA, 1% SDS). Chromatin was sonicated to yield DNA fragments of 200–1000 bp using a Vibra Cell Ultrasonic Processor (Sonics and Materials, INC). The lysate was cleared at 16 100 g for 10 min at 4°C, and samples were diluted with 1.8 ml of ChAP dilution buffer (16.7 mM Tris–Cl, pH 8.1, 167 mM NaCl, 1.2 mM EDTA, 1.1% Triton X-100, 0.01% SDS). Twenty microliters of BSA-blocked Strep-Tactin beads were added to each sample and incubated at 4°C overnight. Beads were washed 5 times with cold RIPA buffer containing 50 mM Tris–Cl, pH 7.5, 1% NP-40, 1% sodium DOC, 0.1% SDS, 1 mM EDTA, 1 M NaCl, 1.5 M Urea, 0.2 mM PMSF, at 15 min for each wash. This was followed by three washes with cold TBS containing 20 mM Tris–Cl, pH 8.0, 150 mM NaCl, 1 mM EDTA, 10 min for each wash. After elution with buffer E (20 mM Tris–Cl, pH 8.0, 150 mM NaCl, 1 mM EDTA, 2.5 mM dithiobiotin), the NaCl concentration was adjusted to 200 mM. Crosslinking was reversed by incubation at 65°C for 8 h, and ChAP DNA samples were recovered by ethanol precipitation in the presence of 20 μg of glycogen and resuspended in 50 μl TE buffer.

Z-DNA library construction

ChAP DNA fragments were blunt-ended by T4 DNA polymerase. Adaptor-L (5′-AAGGAGGAGAGTTATGGATCCCGAC3′) and adaptor-S (5′-CGGTCCGTGATCCGATCAC3′) were annealed and ligated to ChAP DNA fragments using T4 DNA ligase (NEB). Fragments were amplified with adaptor-L for 25 cycles using Taq PCR Core Kit (Qiagen) with 1 × Q solution. PCR products were separated on 0.5% agarose gel in 0.5 x TBE buffer and extracted, followed by digestion with BamHI and separated again on a 1.0% agarose gel. DNA fragments were ligated into the pTZ18R vector cleaved with BamHI and treated with CIP (NEB). DNA was purified by phenol/chloroform extraction and ethanol precipitation, and transformed into 40 μl of Stbl4 competent cells using electroporation (Invitrogen). The resulting DNA library was sequenced at the Beijing Genomics Institute. In total, about 10 500 white colonies were picked and sequenced with primer pTZ-L (5′-GATTAC GAATTTAATCGACTACTA-3′).

ChAP-PCR

Sixteen hotspots consisting of at least four ChAP sequences were subjected for further analysis using ChAP-PCR. PCR primers on the hotspots were designed by Primer3 (http://frodo.wi.mit.edu/cgi-bin/primer3/ primer3 www.cgi) or Vector NTI Suite® 8 (Invitrogen). A new batch of ChAP DNA was used as template and each PCR was optimized individually. PCR products were analyzed on agarose gels after EtBr staining. PCR using genomic DNA as template was used as control to check the quality of primers. In order to confirm the specificity of Z$_{ADAR1}$ binding, seven pairs of primers for hotspot regions or nonhotspot regions were designed, and PCR reactions were performed at 2 mM MgCl$_2$ and $T_a = 60^\circ$C with 35 cycles. PCR products were quantified with Bio-Rad Quantity One software.

RESULTS

We wrote a computer program dubbed Z-catcher in Perl that identifies ZDRs in entire genomes as a function of $\sigma$,.
the DNA superhelical density of protein-free DNA (http://vhp.ntu.edu.sg/zdna/Z_Catcher.zip). Such an approach provides a useful quantitative value that directly indicates the probability for a particular ZDR to be in the left-handed conformation in the human genome and is, therefore, more informative than previous strategies using a pre-set \( \sigma \) value and the program Z-hunt (14,15). Although \( \sigma \) of overall unconstrained (−) supercoiling in the human genome appears null (19), that of transcription-induced, localized (−) supercoiling reaches high levels, even in the presence of eukaryotic topoisomerases (20,21).

When we subjected the human genome sequence to Z-catcher at \( \sigma = -0.07 \), we found that in agreement with previous in silico analyses using Z-hunt, the occurrence of ZDRs is evenly distributed over the set of human chromosomes (data not shown). In order to directly compare ZDR predictions made by both programs, we used two sets of sequences, labeled Demo 1 and Demo 2, which we randomly selected from the human genome as test sequences. Each set is composed of 24 sequences derived from chromosomal positions 8400001 to 8750000 in Demo 1, and from positions 42000001 to 42350000 in Demo 2. Sequences containing undefined bases “N” were excluded, since neither Z-hunt nor Z-catcher can analyze them.

The results of these comparisons show that at \( \sigma = -0.07 \), most ZDRs (91.7%) predicted by Z-catcher were also identified by Z-hunt (Figure 1A). It is also clear that at this \( \sigma \) level, Z-hunt returned more ZDRs than Z-catcher. This changed when we chose \( \sigma = -0.075 \). In this case, Z-catcher returned about 1.5 times more ZDRs than at \( \sigma = -0.07 \). These ZDRs include the majority of those identified by Z-hunt, plus a number of additional ZDRs on each chromosome (Figure 1B). This trend continued when we changed \( \sigma \) to −0.08. Now, the majority of ZDRs returned by Z-catcher is not identified by Z-hunt, while nearly all ZDRs returned by Z-hunt are also predicted by Z-catcher (Figure 1C). Hence, these data show that Z-catcher is a very useful alternative to Z-hunt for in silico ZDR predictions. A detailed description of the former can be found in Supplementary Data.

Next, we used Z-catcher to perform fine mapping of ZDRs over the entire human genome. The results revealed that they begin to cluster at \( \sigma = -0.055 \) between nucleotides +100 and −600 relative to TSS (Figure 1D). Interestingly, a \( \sigma \) level of −0.06 is in the range of that determined for SV40 DNA purified from nuclei of infected mammalian cells (22). Hence, it seemed indeed likely that some of these ZDRs might adopt a Z-conformation in vivo due to localized (−) DNA supercoiling that may result from nucleosome displacement, transcription, or other changes in chromatin structure, as previously reported for the CSF1 promoter (23).

In order to verify that ZDRs predicted by Z-catcher near TSS are in the Z-conformation, we used cell-based assays and ZADAR1 as the probe (24). We generated a recombinant ZADAR1 version containing two short tags fused to the C-terminus (Figure 2A). As control probe which cannot recognize Z-DNA, termed ZADAR1mut, we replaced both tyrosine 177, which specifically interacts with a purine residue in the \( \text{syn} \) conformation, and asparagine 173, which is engaged in water-mediated contacts with the Z-DNA phosphate backbone, with alanine (Figure 2A and B). Previous data revealed that each of these substitutions alone significantly reduced the binding affinity of ZADAR1 to Z-DNA (25).

ZADAR1 and ZADAR1mut were expressed in Escherichia coli and purified to homogeneity. We used pull-down assays and confirmed that ZADAR1 binds specifically to d(GC)\(_6\) in the Z-conformation at 150 mM NaCl (Figure S2). Importantly, only ZADAR1 could be crosslinked with 0.5% formaldehyde to Z-DNA, as demonstrated by EMSA using a genomic target sequence in either the Z- or B-conformation (Figure 2C and D).

Having established binding specificities and conditions for our probes, we used human A549 lung cancer cells to identify genomic DNA segments in the Z-conformation. We employed a ChAP strategy with dual formaldehyde crosslinking. Cells were first fixed through crosslinking directly on culture plates, incubated with purified probes, which was followed by a second round of crosslinking and ChAP (Figure 3A). After ChAP, DNA fragments were released, linked to adaptors, amplified by PCR, and cloned. Analysis of a sample of the amplified ChAP DNA before cloning revealed that a significant amount of template DNA co-purified only with ZADAR1 (Figure 3B and C).

The resulting genomic library contained about 13,000 clones. After sequencing plus data processing which removed adaptor/vector sequences, 10,005 sequences were subjected to BLASTN (http://blast.ncbi.nlm.nih.gov) or Blat (http://genome.ucsc.edu) for alignment with the human genome build 36.2 (NCBI). We found that 7321 sequences were at least 30 bp long and matched with an identity \( \geq 90\% \) and not more than a 1% gap. Total 1715 sequences were mapped to the \( \text{E. coli} \) genome and 969 were not identified in the NCBI nr (nonredundant) database. Lists of the 10,005 sequences used for the alignment and the 969 unidentified sequences can be found at (http://vhp.ntu.edu.sg/zdna/data.htm).

As summarized in a flow chart (Figure 4), we defined a Z-DNA hotspot as a sequence covered by two or more ChAP fragments that either overlapped or are separated by \( \leq 100 \) bp. This definition was based on the following reasoning: First, genomic fragments containing ZDR(s) in Z-conformation recognized by ZADAR1 will inevitably exhibit different lengths due to sonication prior to ChAP. This eventually generates a set of overlapping sequences in our library and is, therefore, a strong indicator for specific ZADAR1 binding. Second, if a genomic region is (−) supercoiled, it is plausible to expect that some ZDRs in Z-conformation might appear in clusters. Therefore, we choose a rather conservative figure of 100 bp as the maximal acceptable distance separating two sequences present in our library. Third, two or more identical ChAP sequences were not considered as they could have been generated as a result of PCR/cloning. It should be highlighted that although we cannot exclude the possibility that sequences which occur only once in our library contain a ZDR(s) in Z-conformation recognizable by our probe, we do not consider them here for further detailed analysis.
A total of 350 hotspots fulfilled these criteria. We eliminated next those hotspots with identical sequences which are located at different genomic regions, since this feature made it impossible to determine which one was actually bound by Z$\alpha$ADAR1. The remaining fell into two distinct categories: those that were defined by ChAP fragments which mapped to a unique region in the genome, labeled as ‘unique hotspots’, and so-called ‘high potential hotspots’. The latter were generated by ChAP fragments that individually were found in multiple sequence repeat regions in the genome; however, they clustered according to our hotspot criteria mentioned above only at one unique genomic region (Figure S3). In total, 122 unique and 64 high potential hotspots were finally mapped to

Figure 1. Predictions of ZDRs by Z-catcher. (A–C) Comparison of ZDRs identified by Z-catcher and Z-hunt. See text for details. Note that the randomly selected sequences from chromosomes 13, 14, 15, 21 and 22 in Demo 1 contained undefined bases, which prevented in silico analyses. The same applied to sequences from chromosomes 16 and Y in Demo 2. (D) Distribution of ZDRs at various $\sigma$ levels around TSS in the human genome. All 103,542 representative TSSs of transcription units in the human genome from the Database of Transcriptional Start sites (DBTSS release 5.2.0) were selected. Flanking regions of TSSs 2000 bp upstream and 2000 bp downstream were extracted and subjected to Z-catcher, with $\sigma$ values ranging from -0.070 to -0.050. The distances between the predicted ZDRs and their corresponding TSSs were calculated and plotted.
We found 66 hotspots in transcribed regions, based on ESTs and mRNAs listed in the UCSC Genome Browser. A closer inspection revealed that the majority (49/74.2%) of these hotspots were located in introns. The remaining fraction was found within exons or splice sites. The level of transcription of these 66 hotspots in A549 cells was checked against the NCBI GEO profile GSM94306 database. Only 31 hotspots matched, with 10 transcribed at high and 21 at low levels.

A previous study found Z-DNA formation in discrete regions of the human c-myc gene in the vicinity of TSSs. The structural transitions there seemed to be linked to transcription in permeabilized nuclei (26). Since c-myc is transcribed in A549 cells, yet no ZDR of this region is present in our hotspot list, we employed ChAP–PCR and the same primer pairs used earlier (26) to determine whether we can detect an enrichment of these c-myc regions in the Z_{ADAR1} sample when compared with the control sample obtained with Z_{ADAR1}mut. However, the results showed that no enrichment is seen (data not shown).

We found that 136 hotspots belonged to tandem repeat families; 49 are ALR/alpha satellite sequences and 22 were found in Alu subfamilies. The rest belonged to different repeat classes, including LTR/L1 and MER. Interestingly, 34 hotspots in the ALR/alpha satellite family and 12 non-alpha satellite hotspots mapped to centromeres on 13 different chromosomes (Figure 5). An analysis via the RepeatMasker Web Server (http://www.repeatmasker.org/) identified six non-alpha satellite hotspots as HSATII satellites, one as SST1 satellite, two as BSR/beta satelltes and three as nonrepeat sequences. Of the 46 centromeric hotspots, 35 were high potential Z-DNA hotspots, and an analysis of all centromeric hotspots via Z-catcher revealed that 36 were predicted to flip into Z-DNA at a (–) σ level between 0.07 and 0.09, while the remaining 10 required higher levels of torsional strain. However, this was not a particular feature of centromeric hotspots as the remaining 140 also showed a similar dependence for high levels of supercoiling (predicted ZDR sequences within hotspots are listed in Table S4). In addition, the 6887 nonhotspot sequences in our library showed the same tendency to undergo structural transitions at (–) σ levels equal to or below 0.08 (data not shown).

In order to provide further evidence that our map of Z-DNA segments was made up of specific binding sites for Z_{ADAR1}, ChAP–PCR was employed using a different biological sample as starting material. We chose 33 hotspots, each defined by at least four ChAP sequences in our library. We were able to design specific primers for 16 hotspots (Table S5). The results showed that, with two exceptions, Z_{ADAR1} ChAP samples were markedly enriched with hotspot fragments when compared with Z_{ADAR1}mut or BSA control samples (Figure S4A).

We asked next whether these fragments were also enriched over random fragments in the Z_{ADAR1} ChAP sample. A second round of ChAP–PCR was therefore performed and normalized to control PCRs using genomic DNA as templates. We designed seven pairs of primers for both hotspot and nonhotspot regions (Table S6).
The signal of each PCR was quantified and ratios between ChAP–PCR and control PCRs were calculated and plotted. The results showed that six hotspot PCRs produced stronger signals than nonhotspot PCRs (Figure S4B and C).

The syn conformation of purines in Z-DNA is more accessible to solvent (27), and the two extruded nucleotides at B–Z junction are potential targets for chemical modification (28). This led us to investigate whether a correlation existed between Z-DNA hotspots and the occurrence of single nucleotide polymorphisms (SNPs) listed in NCBI dbSNP build 128. Both validated RefSNPs and nonvalidated RefSNPs, i.e. all entries in the database, were used for this analysis. RefSNPs in hotspots and in three sections of flanking regions were counted and displayed as SNP density (Figure 6A). Pair-wised Wilcoxon Signed-Rank test was used to compare SNP densities in hotspots and in flanking regions. Interestingly, the 46 hotspots in centromeres correlated with the occurrence of SNPs when validated RefSNPs were considered (Figure 6B, Table 1). This correlation was even more significant ($P < 0.001$) with nonvalidated RefSNPs. When we considered only the 11 unique hotspots in centromeres with nonvalidated RefSNPs, this correlation was still significant ($P < 0.05$). However, hotspots found outside centromeres ($n = 140$) did not reveal a significant enrichment of SNPs ($P > 0.1$) (Table 1).

**DISCUSSION**

Our study provided a first Z-DNA map of the human genome generated by a Z-DNA specific, cross-linkable protein probe, termed Z$_{ADAR1}$. The novel ChAP strategy employed here was based on a previous finding that formaldehyde treatment does not affect binding of Z$_{ADAR1}$ to Z-DNA (10). It was shown that d(TG) repeats in the CSF1 promoter region were cleaved by Z$_{ADAR1}$FOK in cells fixed with 1% formaldehyde. Z$_{ADAR1}$FOK is a recombinant restriction enzyme created by the fusion of the nuclease domain from FokI endonuclease with two tandem copies of Z$_{ADAR1}$ (29).

In our protocol, A549 cells were treated in a similar way and incubated with our protein probes before the second crosslinking, followed by ChAP. Based on the following observations and reasoning, we think that our approach is unlikely to induce B-to-Z structural transitions in
appropriate sequence stretches. First, DNA sequences retrieved from ChAP contained very few d(GT)$_n$ or d(GC)$_n$ repeats. Because d(GT)-containing microsatellites are abundant in the human genome and expected to flip into Z-conformation at moderate levels of (−) supercoiling, it seems unlikely that many of our hotspots were generated by, for example, nucleosome displacement as a result of the chosen experimental strategy. Second, confirmation of our results by ChAP–PCR using a different biological sample indicated that crosslinking of Z$_\text{ZADAR1}$ occurred at specific genomic DNA segments. Third, the fact that cells were crosslinked before binding of Z$_\text{ZADAR1}$ made it unlikely that the probe induced a B-to-Z transition, as observed with oligonucleotides in vitro (24). We think that the accompanying introduction of positive supercoiling will impose a very strong energetic barrier in a fixed chromatin background, which prohibits efficient diffusion of supercoils along the chromatin fiber, unless a chromatin domain is already under negative torsional strain that is insufficient to drive a B-to-Z transition. In the latter case, the introduction of positive supercoils due to binding of Z$_\text{ZADAR1}$ would be cancelled.

Having established a first Z-DNA map of the human genome by applying very stringent selection criteria, a main conclusion is that the in silico predicted enrichment of ZDRs in Z-conformation near TTS could not be verified. In fact, only two hotspots were found near TSS. This finding does not exclude, however, a possibility that some of these ZDRs adopt Z-conformations in other human cells, or undergo structural transitions as a result of different metabolic states of A549 cells. Furthermore, it remains possible that conformation-specific DNA binding proteins mask some Z-DNA sequences in promoter regions. However, the fact that the A549 chromatin is fixed through histone–DNA crosslinking, which also freezes DNA in its topological state, indicates that a pure thermodynamic-driven in silico approach to predict Z-DNA segments in a genome may have severe limitations. The kinetics of Z-DNA formation in regions near TSS may be too slow, perhaps due to efficient diffusion of transcription-induced (−) supercoiling in these regions. Also, the occupancy of ZDRs near TSS by transcription factors could stabilize the B-conformation. Although some ZDRs in promoter regions may actually adopt the

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**Figure 5.** Z-DNA hotspots mapped to human chromosomes. One hundred and eighty six hotspots were identified and classified as those located in transcribed regions, in centromeres, and unspecified genomic regions. Hotspots were mapped on chromosomes using UCSC Genome Browser, and chromosomes are drawn to scale. Detailed information on all hotspots is listed in Table S3.
Z-conformation in vivo, as recently shown for the CSF1 gene (10), our data indicate that Z-DNA formation is unlikely to be a key component of a widespread mechanism regulating transcription initiation.

We found that 66 hotspots are located in transcribed genomic regions, and 49 of them were in introns. The formation of Z-DNA there might be linked to transcription-induced supercoiling, as suggested previously for the c-myc gene, corticotropin-releasing hormone gene and beta-globin gene (26,30,31). However, we could not confirm Z-DNA formation in the c-myc gene, which might, at least in part, be due to different protocols employed for binding of protein probes to chromatin. Furthermore, the use of a database of transcriptional profiling confirmed that 31 of the 66 hotspots are actually transcribed in A549 cells, with 21 transcribed at low and 10 at high level. No data could be found in the database for the remaining 35 hotspots, which leaves the possibility open that these are also transcribed at some level. We conclude, therefore, that no strong correlation existed between the level of transcription and Z-DNA formation.

A striking result of our study is that 46 (25%) hotspots were located in centromeres of 13 chromosomes. Computational analysis using Z-catcher revealed that all of them will show a structural transition into Z-DNA at \( \sigma < -0.09 \), which is a rather high level of unconstrained (–) supercoiling. It is known that functional eukaryotic centromeres have an irregular nucleosome positioning (32). This may contribute to the generation of high \( \sigma \) levels, perhaps due to active chromatin remodeling. Formation of Z-DNA may in fact stabilize this situation since Z-DNA cannot be incorporated into nucleosomes (8,9). In addition, histone H3 variant CENP-A is found in centromeres and contributes there to a more rigid nucleosome conformation (33). This might also aid in the build-up of torsional strain. It has been reported that scaffold/matrix attachment regions are more abundant in centromeres or neocentromeres than in chromosome arm regions (34). It is very likely that they demarcate regions under high torsional strain and prevent diffusion of negative supercoiling into neighboring chromatin domains. Finally, the prominence of multiple topoisomerase II cleavage sites in human centromeres could imply that these regions have a high level of unconstrained supercoiling (35).

Centromeres are spindle attachment sites during mitosis and meiosis, and they mediate proper segregation of chromatids into daughter cells. However, centromeres are extremely diverse in sequence composition, ranging from the 125 bp so-called point sequence in Saccharomyces cerevisiae to highly repetitive satellite DNA in vertebrates. In fact, centromere repeats are the most rapidly evolving sequences in eukaryotic genomes (36), involving sequence expansions and contractions. It has been proposed (36) that these repeat variations might provide a functional advantage to centromeres during female asymmetric meiosis. Based on our results, it is possible that some of these gross deletions plus gene conversions might be triggered by formation of Z-DNA, which was shown to induce genomic recombination (11,37,38).

Alpha-satellites are highly variable in sequence composition (39). Careful examination of SNP distributions in centromeres revealed that they are not evenly spread but clustered in certain segments. Our finding that high SNP densities correlated with high densities of SNPs.

Table 1. P values of Wilcoxon Signed-Rank tests of SNP densities

| Flanking regions | 0.5 kb | 1 kb | 2 kb |
|------------------|--------|------|------|
| All_RefSNPs      | <0.001 | <0.001 | <0.001 |
| Validated_RefSNPs| 0.014  | 0.045 | 0.042 |
| All_RefSNPs      | 0.021  | 0.026 | 0.013 |
| Validated_RefSNPs| 0.075  | 0.041 | 0.05  |
| All_RefSNPs      | 0.283  | 0.211 | 0.388 |
| Validated_RefSNPs| 0.865  | 0.816 | 0.942 |
| All_RefSNPs      | 0.11   | 0.103 | 0.089 |
| Validated_RefSNPs| 0.238  | 0.325 | 0.219 |

Distributions of RefSNPs were normalized as SNP densities (SNP/bp) for statistical analysis, which showed that only hotspots in centromeres correlated with high densities of SNPs.
anchor points for the kinetochore during mitosis and meiosis (40), some of their structural features related to function should be conserved. This could lead to Z-DNA formation in neighboring sequences, which might be functionally less constrained and more prone to be mutated and to accumulate SNPs over generations. Z-DNA formation could then serve there as a buffer against very high levels of (−) supercoiling which may build up during mitosis.

Our study provided a strategy to generate a first snapshot map of the most stable Z-DNA segments in the human genome. It will be interesting to compare this map in the future with those obtained during different stages of the cell cycle or with other cell types. Since the formation of Z-DNA can be regarded as a real time indicator for genetic activities in a genome (3), comparative studies could ultimately lead to a better understanding of genome-wide chromatin dynamics and genetic stability.

SUPPLEMENTARY DATA
Supplementary Data are available at NAR Online.

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