The transcriptional co-activator LEDGF/p75 displays a dynamic scan-and-lock mechanism for chromatin tethering

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Received May 2, 2010; Revised September 27, 2010; Accepted September 28, 2010

ABSTRACT

Nearly all cellular and disease related functions of the transcriptional co-activator lens epithelium-derived growth factor (LEDGF/p75) involve tethering of interaction partners to chromatin via its conserved integrase binding domain (IBD), but little is known about the mechanism of in vivo chromatin binding and tethering. In this work we studied LEDGF/p75 in real-time in living HeLa cells combining different quantitative fluorescence techniques: spot fluorescence recovery after photobleaching (sFRAP) and half-nucleus fluorescence recovery after photobleaching (hnFRAP), continuous photobleaching, fluorescence correlation spectroscopy (FCS) and an improved FCS method to study diffusion dependence of chromatin binding, tunable focus FCS. LEDGF/p75 moves about in nuclei of living cells in a chromatin hopping/scanning mode typical for transcription factors. The PWWP domain of LEDGF/p75 is necessary, but not sufficient for in vivo chromatin binding. After interaction with HIV-1 integrase via its IBD, a general protein–protein interaction motif, kinetics of LEDGF/p75 shift to 75-fold larger affinity for chromatin. The PWWP is crucial for locking the complex on chromatin. We propose a scan-and-lock model for LEDGF/p75, unifying paradoxical notions of transcriptional co-activation and lentiviral integration targeting.

INTRODUCTION

Gene expression is regulated by transcriptional cofactors that fine-tune the interaction of the general transcription machinery with gene-specific transcription factors. Lens epithelium-derived growth factor (LEDGF/p75) was originally identified as a 75-kDa transcriptional co-activator interacting with the VP16 activation domain and with components of the general transcription machinery (1). LEDGF/p75 (530 amino acids) shares the first 325 amino acids with p52, an alternative splice variant from the same PSIP1 gene (1,2) (Figure 1A) and plays an important role in cell survival (3), oncogenesis (4–7), autoimmunity (8,9) and integration and replication of the human immunodeficiency virus type 1 (HIV-1) (10,11).

LEDGF/p75 has an extensive interactome (Figure 1B). The protein contains multiple DNA/chromatin binding domains and a conserved protein–protein interaction domain. The N-terminal PWPP domain of LEDGF/p75 (amino acids 1–93) holds a conserved (though not invariant) Pro-Trp-Trp-Pro motif and belongs to the Tudor domain ‘Royal Family’ of protein domains regulating the chromatin function (12,13). This domain is generally involved in chromatin structure regulation through protein–protein interactions (14). A tripartite element in LEDGF/p75 consisting of the two AT-hooks (amino acids 191–197 and amino acids 178–183) and the nuclear localization signal (NLS) (amino acids 148–156) cooperates with the PWWP domain for interaction with DNA/chromatin, as has been shown in vitro (15) and in vivo (16). LEDGF/p75 allegedly interacts with heat shock elements (HSE) through its helix-turn-helix (HTH) motifs (amino acids 421–442 and amino acids 471–492) and with...
stress-related regulatory elements (STRE) through the PWWP domain to specifically promote expression of stress-related genes (17–21). LEDGF/p75 contains a high percentage of charged residues (39.4%) and contains two defined regions with positively charged residues (amino acids 94–142 and amino acids 208–325), that are involved in electrostatic interactions with DNA/chromatin (15). A conserved superhelical domain in LEDGF/p75 was originally identified as the domain for binding to HIV-1 integrase (IN), the enzyme that catalyzes the integration of HIV in the genome of an infected cell (22,23). By virtue of this interaction LEDGF/p75 biases lentiviral integration towards actively transcribed regions in the genome (24–26). The 'integrase binding domain' (IBD) of LEDGF/p75 (amino acids 347–429) shows structural homology to known protein–protein interaction motifs (27) and has been shown to interact with multiple other proteins in cells: (i) JPO2, a Myc transcription factor interacting protein (28), (ii) the menin tumor suppressor, implicated in cancer and transcriptional regulation as a component of the MLL-HMT (mixed lineage histone methyltransferase) complex (7), (iii) PogZ, pogo transposable element derived protein with a Zinc finger, a domesticated transposase (29) and (iv) heterodimeric Cdc7/ASK (30). Cdc7 is a Ser/Thr kinase essential for the initiation of DNA replication throughout the S-phase and its activity is controlled via interaction with a regulatory subunit, activator of S-phase kinase (ASK).

In this work we study the intracellular mobility of LEDGF/p75, in particular chromatin interactions, quantitatively and in vivo with a combination of fluorescence techniques. We label LEDGF/p75 with eGFP and perform a detailed characterization of the hybrid protein. We provide new insight in the transcription factor chromatin binding kinetics in vivo. We investigate interaction partner mediated chromatin tethering of LEDGF/p75 and demonstrate the role of the PWWP domain for this.

MATERIALS AND METHODS

Cell lines
HeLa cells were obtained from the NIH Reagent program and were grown in ‘Complete Medium’, high-glucose...
Dulbecco’s Modified Eagle Medium (Gibco BRL, Belgium), supplemented with 10% heat-inactivated fetal bovine serum (Sigma-Aldrich gmbh, Taukirchen, Germany) and 50 μg/ml Gentamyce (Gibco BRL, Belgium), at 5% CO2 and 37°C in a humidified atmosphere. The generation of monoclonal HeLa cells stably suppressing endogenous LEDGF/p75 (HeLa-p75KD) has been described earlier (31). An average HeLa cell nuclei volume V = \( \frac{4}{3} \pi \left( \frac{W}{2} \right)^{3} \) was calculated by measuring the length (L) and width (W) of seven nuclei by optical transmission microscopy and assuming an ellipsoidal nuclear shape. Assuming 10–100,000 genes per cell, this roughly corresponds to a gene concentration of \( \sim 5–100 \) nM.

Western blotting, cellular fractionation assays

Western blotting was performed on whole cell lysates as described before (32) with a specific anti-LEDGF/p75 antibody (A300-847A, Bethyl Laboratories, Montgomery, TX, USA). Cellular fractionation assays were performed as described earlier (15), with the specific LEDGF/p75 antibody, an anti-RFP antibody for mRFP-IN (AB3216, Millipore N.V., Brussels, Belgium), an in-house polyclonal anti-GFP antibody and anti-α-tubulin (T5168, Sigma-Aldrich, Bornem, Belgium). Briefly, 24 h post-transfection cells were harvested and either lysed for checking overall expression (T) or extracted with Triton X-100 to separate non-chromatin binding fraction (P2). DNasel-treatment allowed subsequent separation into a DNaseI-treatment allowed subsequent separation into a non-chromatin binding fraction (P2).

Transfections

For the transient transfections 0.5–1 \times 10^5 cells were seeded per well in a Lab-Tek™ Chambered Coverglass (VWR International, Leuven, Belgium) to obtain 50–70% cell confluency after overnight incubation. Transfections were performed with a Mirus TransIT™-HeLaMONSTER™ transfection kit (VWR International, Leuven, Belgium), with 1 μl TransIT reagent, a maximum of 0.5 μg plasmid DNA per well and 1 μl MONSTER reagent per μg of DNA. After the transfection mixture was prepared in 50 μl fresh OptiMEM per well (Gibco BRL, Belgium), 450 μl prewarmed Complete Medium was added and this mixture was slowly added to the cells and the cells were incubated for at least 12 h.

Theory: quantifying interactions with immobile structures

Consider a binding equilibrium of a protein L that can interact with an immobilizing binding site S on the chromatin:

\[
\text{L+S} \stackrel{K_{\text{ass}}}{\longrightarrow} \text{LS} \quad \text{with} \quad K_{\text{ass}} = \frac{[\text{LS}]}{[\text{L}][\text{S}]} \quad (1)
\]

Association rate = \( k_{\text{on}}[\text{L}][\text{S}] \quad \text{slow} \rightarrow (k_{\text{on}}S_{0})[\text{L}] = k_{\text{on}}^{*}[\text{L}] \quad (2) \)

Dissociation rate = \( k_{\text{off}}[\text{LS}] \quad (3) \)

\( t_{\text{free}} = 1/k_{\text{on}}^{*} \) and \( t_{\text{bound}} = 1/k_{\text{off}} \quad (4) \)

with [L] the concentration of free protein L, in this case, LEDGF/p75, [S] the concentration of free immobile binding sites, [LS] the concentration of protein L bound to a binding site, \( K_{\text{ass}} \) the association binding constant, \( k_{\text{on}}^{*} \) the rate constant for association and \( k_{\text{off}}^{*} \) the rate constant for dissociation. When the concentration of binding sites is high, \( k_{\text{on}}^{*} = k_{\text{on}}S_{0} \) is the pseudo first order rate constant for association. The \( t_{\text{free}} \) is the average time a protein spends between two successive binding events. The \( t_{\text{bound}} \) is the residence time on the chromatin binding site. Intracellular binding reactions with immobile structures (nuclear envelope, chromatin, membrane) can be studied with fluorescence recovery after photobleaching (FRAP) (33), continuous photobleaching (CP) (34,35) and fluorescence correlation spectroscopy (FCS) (36). Generally, four dynamic regimes can be thought of:

Fast association (\( t_{\text{free}} \leq t_{\text{diff,free}} \)) and slow dissociation (\( t_{\text{bound}} > t_{\text{diff,free}} \)). The protein binds with high affinity. It will be photobleached in the laser focus when FCS is performed. In this case, FRAP is a good method to analyse the binding kinetics: the dynamics will be slow and a model for reaction and diffusion is necessary because of the high probability of association (33).

Slow association (\( t_{\text{free}} > t_{\text{diff,free}} \)) and very slow dissociation (\( t_{\text{bound}} > 100 \text{ ms} \)). The chance of binding is low but once bound the fluorochrome will be photobleached when measured with FCS. The fraction of the signal (from CP or FCS) that photobleaches is a measure for the \( K_{\text{ass}} = K_{\text{ass}}[\text{sites}] \) (36):

\[
\frac{F_{\text{bleached}}}{F_{\text{unbleached}}} = \frac{[\text{LS}]}{[\text{L}]} = \frac{k_{\text{on}}^{*}}{k_{\text{off}}^{*}} = K_{\text{ass}} \quad (5)
\]

When Equation 1 is solved to [LS] in function of total concentrations \( L_{0} \) and \( S_{0} \), \( K_{\text{ass}} \) (\( K_{\text{diff}} \)) and the concentration of binding sites can be determined by non-linear least-squares fitting of the percentage photobleaching versus the total concentration of protein \( L_{0} \):

\[
F_{\text{bleached}} = \left( L_{0} + S_{0} + K_{\text{diff}} \right) - \sqrt{\left( L_{0} + S_{0} + K_{\text{diff}} \right)^{2} - 4L_{0}S_{0}} \quad (6)
\]

The total concentration \( L_{0} \) can be measured with FCS. The parameter \( y_{0} \) is a constant accounting for the constant fraction of photobleaching at high concentrations. Alternatively, diffusion independent FRAP can be used to study the binding reaction (33).

Fast association (\( t_{\text{free}} < t_{\text{diff,free}} \)) and fast dissociation (\( t_{\text{bound}} < t_{\text{diff,free}} \)). The binding reaction is observed as a diffusion component in FRAP and FCS. The ‘effective’ slow diffusion coefficient defines the affinity (33,36,37):

\[
D_{\text{observed}} = \frac{D_{\text{free}}}{1 + k_{\text{on}}^{*}k_{\text{off}}^{*}} = \frac{D_{\text{free}}}{1 + t_{\text{bound}}/t_{\text{free}}} = D_{\text{free}}F_{\text{free}} \quad (7)
\]

In this regime, only the ratio of rate constants can be determined from the ACF. The diffusion coefficient of
free cellular LEDGF/p75 was calculated with (assuming globular proteins):

$$D_{\text{free eGFP-LEDGF/p75}} = D_{\text{eGFP}} \sqrt{\frac{M_r \text{eGFP}}{M_r \text{eGFP-LEDGF/p75}}}$$

(8)

$M_r$ being the relative molecular mass and $D_{\text{eGFP}} = 33.0 \mu m^2/s$ (as measured with FCS).

**Slow association** ($t_{\text{free}} >> t_{\text{diff}}$), moderate dissociation ($\tau_{\text{diff,free}} < \tau_{\text{bound}} < 100 ms$). The binding reaction is observed as an exponential component in FRAP and FCS (33,36). The fraction of free and ‘apparent slow’ diffusion components in FCS is a measure for the affinity:

$$\frac{F_{\text{slow}}}{F_{\text{fast}}} = \frac{[\text{S}]}{[\text{L}]} = \frac{k_{\text{on}}}{k_{\text{off}}} = K_{\text{ass}}$$

(9)

In conclusion, in the first two regimes the binding reaction is not and in the last two regimes the binding reaction is observed during an FCS experiment.

**Spot fluorescence recovery after photobleaching** and **half-nuclear fluorescence recovery after photobleaching**

Diffusion rates and binding reactions of cellular proteins with slow/immobile cellular structures can be quantified with spot fluorescence recovery after photobleaching (sFRAP) (33,38). By illuminating a defined region-of-interest in the sample with a brief high-intensity laser pulse, the fluorescence is rapidly photobleached. It will recover due to the exchange of bleached with unbleached molecules. FRAP measurements were performed on a LSM510/ConfoCor2 (Carl Zeiss, Jena, Germany) at 488-nm excitation. Since bleaching-while-acquisitioning (39) and recovery-while-bleaching (40) influence the shape of the FRAP curve, careful optimization of laser power and imaging parameters was necessary.

sFRAP measurements were performed in a 1-μm radius (ω) circular spot. For eGFP and eGFP-LEDGF/p75, acquisitioning was every 65 ms at 5 μW excitation power (front objective aperture) and bleaching was performed with a single 65-ms scan at 1.6 mW. For eGFP-LEDGF/p75 in the presence of mRFP-IN, acquisitioning was every 5 s and bleaching was performed with a single 170-ms scan. Spot FRAP data represent averages of at least 10 cells. Analysis was performed with a standard diffusion model (38), a reaction-dominant model (33) or a full diffusion-reaction model (33), as described in detail in the Supplementary Information.

Half-nuclear FRAP (hnFRAP) measurements (41,42) were performed by photobleaching half of the nucleus with a single scan (1.4 s, 1.6 mW) and monitoring the subsequent fluorescence redistribution (every 1.4 s, 5 μW). Average nuclear pixel intensities per horizontal image line were calculated for each image with an ImageJ plug-in. Next, each post-bleach data point was normalized by the total average cell intensity, to account for whole-cell photobleaching. Next, each normalized data point was normalized by the corresponding pre-bleach data point, to account for cellular fluorescence inhomogeneities. Finally, vertical pixel distance was converted to an absolute distance scale (1 pixel = 90 nm) and data were plotted in a graph and fitted with (43):

$$F_{\text{norm}}(x) = \frac{1 - F_{\text{bleach}}}{2} \left[ \text{erf} \left( \frac{b - x}{\sigma(t)} \right) \right]$$

(10)

with $F_{\text{norm}}(x)$ the normalized fluorescence intensity as a function of the position $x$ (μm), $F_{\text{bleach}}$ the bleach depth, erf an error function describing the shape of the bleach profile, with $b$ (μm) the position of the bleach border and $\sigma(t)$ the displacement of the proteins as a function of time $t$ (s). HnFRAP data for a single cell were fitted globally with $F_{\text{bleach}}$ as a global fit parameter. From the slope of the mean square displacement $\sigma^2(t)$ the diffusion coefficient $D$ (μm$^2$/s) was calculated (44):

$$\sigma^2(t) = 4Dt$$

(11)

Time zero was approximated with the moment during acquisition where half a bleaching iteration had been performed. HnFRAP data represent averages of at least five cells.

**FCS and fluorescence cross-correlation spectroscopy**

With FCS, diffusion and binding reactions of fluorescently labeled molecules can be studied in solution and in cells (45,46). On a confocal microscope the fluorescence in the sub-micrometer (<10$^{-15}$l) sized confocal volume is measured and then autocorrelated. The low time resolution decaying fluorescence trace contains information about diffusion and binding reactions, the high time resolution autocorrelation function (ACF) in addition provides the absolute protein concentration. For normal FCS fluorescence cross-correlation spectroscopy (FCCS) measurements a commercial FCS/FCCS microscope (LSM510/ConfoCor2, Carl Zeiss, Jena, Germany) was used. Enhanced GFP (47) was excited at 488 nm (3 μW) and mRFP1 (48) was excited at 543 nm (9 μW). The excitation light was reflected by a dichroic mirror (HFT 488/543) and focused through a C-Apochromat 40×/1.2 W Korr/0.13–0.17 objective. The fluorescence emission light was split by a second dichroic mirror (NFT 570) into two separate beam paths and passed through a 505–530 nm bandpass filter and 70-μm pinhole for eGFP fluorescence and a 600–650 nm bandpass filter and 78-μm pinhole for mRFP1 fluorescence. The radial radius of the confocal volume at 1/e$^2$ times its maximal intensity, $\omega_1$, was determined by measuring the $\tau_{\text{diff}}$ of rhodamine 6G in water ($D = 426 \mu m^2/s$) (49). In each cell cytoplasm or nucleus, 10 consecutive fluorescence intensity (detected photons per second) and correlation measurements of 20 s were performed. Correlation functions were fitted with a one- or two-component model, a normal or anomalous diffusion model (50), a reaction-dominant model (36) or a confined diffusion model (43), as described in detail in the Supplementary Information.

**CP**

CP traces were also recorded on the FCS set-up. Individual CP measurements were normalized to the first
Tunable focus FCS

Tunable focus FCS (TFFCS) was performed on a homebuilt set-up (52). The 30-mW 488-nm line of an Ar–Kr laser (Newport Spectra-Physics, Utrecht, The Netherlands) was split 50/50. One beam was maximally focused at the back-aperture of the objective (Olympus UPLFLN 100×/NA1.3/Oil, Olympus Belgium N.V., Aartselaar, Belgium) to create wide-field excitation for observing the fluorescence of the cells through the eyepiece of the microscope (Olympus IX). The other beam was carefully expanded, collimated and directed centrally through an adjustable diaphragm, to allow for TFFCS measurements. A fixed 50-μm pinhole was used for confocal detection on an avalanche photodiode (SPCM-AQR-15, Perkin-Elmer, Wiesbaden, Germany). The excitation power at each diaphragm setting was attenuated to give a constant power/area in the focal spot suitable for cellular FCS, avoiding potential intensity-dependent artefacts. Reproducible switching was achieved by measuring the residual power behind the diaphragm. TFFCS measurements were analysed with a 2-component model for normal diffusion.

In vivo eGFP-LEDGF/p75 exists in a chromatin bound state, a slow state and a fast state

We performed sFRAP experiments in HeLa cells. The fluorescence of the eGFP control exhibited fast and complete recovery, as expected for a free diffusing, inert protein (Figure 2A black). The fluorescence recovery of eGFP-LEDGF/p75 was also complete but much slower (Figure 2A, magenta). A pure diffusion model provided a good fit of the experimental data and a diffusion coefficient $D = 0.41 \pm 0.06 \mu m^2/s$ was calculated (Table 1), which is 53-fold lower than expected for free diffusion (Figure 2A, green, with D from Equation 8). Of concern, while diffusion time of eGFP ($t_{diff} = 97 ms$) could not be determined correctly due to recovery-while-photobleaching (40), the dynamics of eGFP-LEDGF/p75 ($t_{diff} = 2.13 s$) were slow enough to be measured accurately. A reaction-dominant model did not fit the data well and a full diffusion-reaction model did not provide a better fit than a pure diffusion model (Supplementary Figure S1A–C).

Next, we applied CP (39). The CP signal from eGFP in HeLa cells decayed slowly and biexponentially (Figure 2B, black), as has been described before for eGFP (35). For eGFP-LEDGF/p75 (Figure 2B, magenta) a biphasic decay was also observed (Figure 2B, black). Resuming the CP measurement after a pause resulted in a similar decay (data not shown), arguing for the absence of an immobile fraction. It has been shown that slow diffusion in a finite compartment can indeed lead to a biphasic fluorescence decay (34). Settling of diffusion of eGFP-LEDGF/p75 ($D = 0.41 \mu m^2/s$) in a HeLa nucleus ($R = 8.74 \mu m$) should take ~47 s (Supplementary Information), which agrees quite well with the completion of the fast exponential decay we observed (Figure 2B, dashed line). Of concern, fitting the same data with a binding model with medium dissociation rate (Figure 2B, green) did not result in a better description of the experimental data than a simple biexponential decay (35), proving that diffusion and binding are necessary to describe the observed photobleaching.
Interestingly, in the <500 nM concentration range the fraction of the fast exponential was concentration dependent (Figure 2C), which suggests the presence of a low concentration of higher affinity binding sites. In line with this, a full reaction-diffusion model for analyzing the sFRAP data also suggested a low fraction of higher affinity binding sites (Supplementary Table S1). With Equation 6 the data from Figure 2C could be fitted to determine an average concentration of binding sites $S_0 = 61 \pm 31$ nM and a steady state dissociation constant $K_{diss} = 103 \pm 66$ nM.

The fluorescence signal after the initial photobleaching was divided in 20-s intervals which were autocorrelated individually (Figure 2B-top) and an average ACF was calculated with FCS (56). The eGFP exhibited free diffusion in the nucleus (Figure 2D, black and Table 1) with $D = 33.0 \pm 3.5 \mu m^2/s$ (Table 1), which is 2.9 times slower than measured in solution (49) and in good agreement with previous findings in cells if the correct $D$ of the calibration probe ($426 \mu m^2/s$ versus $280 \mu m^2/s$) is taken into account (57). The eGFP-LEDGF/p75 ACF was again strongly shifted into a slower time scale (Figure 2D, magenta) with respect to the calculated ACF for free diffusion (Figure 2D, green). A one-component (normal or anomalous) diffusion model did not describe the ACF well (Supplementary Figure S2A and B). A two-component normal diffusion model described the ACF very well (Supplementary Figure S2C): a fast component (35 ± 3%) with apparent $D = 22.3 \pm 4.1 \mu m^2/s$ and a slow component (65 ± 3%) with $D = 0.5 \pm 0.1 \mu m^2/s$ was determined (Table 1). The slow $D$ is in good accordance with the $D$ obtained from FRAP. A

Figure 2. Dynamic subpopulations of LEDGF/p75 revealed by sFRAP, CP and FCS. (A) sFRAP experiment of eGFP and eGFP-LEDGF/p75 in living HeLa cells. The diffusion coefficient we calculated for eGFP after fitting with a pure diffusion model was only 10.28 $\mu m^2/s$ (Table 1). For eGFP the observed recovery was lower than expected (solid orange curve) because of unavoidable recovery-while-photobleaching, which occurred because the diffusion time of eGFP ($t_{diff} = 97 ms$) was too small with respect to the acquisition time of the microscope (65 ms per iteration on our setup) (40). For eGFP-LEDGF/p75, the arrow indicates the decrease in mobility of eGFP-LEDGF/p75 due to chromatin interactions. $D_{eGFP} = 33 \mu m^2/s$ is the $D$ we measured with FCS, $D_{eGFP-LEDGF/p75} = 22 \mu m^2/s$ is calculated from $D_{eGFP}$ with Equation 8. See also Supplementary Figure S1. (B) CP experiment of eGFP and eGFP-LEDGF/p75. Upper panel is a 80–100-s zoom. (C) Concentration dependence of the amount of photobleaching of eGFP-LEDGF/p75 during the first 20 s. Solid line = fit with Equation 6. (D) FCS experiment of eGFP and eGFP-LEDGF/p75. Solid lines = two-component normal diffusion model. See also Supplementary Figure S2. Error bars = SD.
Table 1. Dynamics of eGFP and eGFP-LEDGF/p75 measured with sFRAP, FCS, CP, TFFCS and hnFRAP

|                | eGFP          | eGFP-LEDGF/p75 |
|----------------|---------------|----------------|
|                | sFRAP | FCS/CP | TFFCS | sFRAP | FCS/CP | hnFRAP | TFFCS |
| Fimm (%)       | 0.1 ± 0.1     | –       | –     | –1.2  | –       | –      | –     |
| Dfree or DFRAP (μm²/s) | 10.28 ± 0.63³ | 33.0 ± 3.5 | 36.4 ± 3.1 | –     | 22.3 ± 4.1¹ | –     | 35.3 ± 10.7 |
| Ffast (%)      | –       | 97.3 ± 0.7 | –     | –     | 35.9 ± 3.0⁰ | –     | 44.1 ± 5.7⁴ |
| Dfast or DFRAP (μm²/s) | –     | 0.8 ± 0.7 | –     | 0.41 ± 0.01 | 0.5 ± 0.1 | 1.02 ± 0.01 | 0.95 ± 0.01 |
| [sites]high (nM) | –     | –     | –     | 1.8 ± 3.1 | 2.2 ± 0.4 | 4.6 ± 0.94 | 4.3 ± 1.0 |
| Kdiss (nM)     | –     | –     | –     | 103 ± 66 | –       | –      | –     |
| [sites]high (nM) | –     | –     | –     | 61 ± 31 | –       | –      | –     |

Values represent averages of at least 10 (sFRAP), 20 (FCS/CP), 10 (TFFCS) or 5 (hnFRAP) measurements and are ±SD.

¹This value is an underestimation, because the shortest bleach iteration (65 ms) was still too long to avoid recovery-while-photobleaching.

²This parameter likely does not represent a diffusion process, see Figure 3F and ‘Discussion’ section.

³The contribution of each component did not depend significantly on the concentration.

⁴The fractional contribution of each component did not depend on the focus spot size.

⁵This parameter was obtained by fitting with a one-component anomalous diffusion fit.

⁶Fit not satisfying, see Supplementary Figure S2B.

⁷Ffree is calculated from the apparent diffusion law, see ‘Materials and Methods’ section, Equation 7, with DeGFP-LEDGF/p75 = 22.3 μm²/s.

The slow moving fraction of LEDGF/p75 (D = 0.5 μm²/s, τdiff = 26 ms) is too fast to represent chromatin controlled movement of associated LEDGF/p75, which is reported to be 2–3 orders-of-magnitude slower (58). We performed TFFCS, where the excitation volume can be tuned in size, to verify the scaling of the observed dynamics with the radius of the measurement spot (33,59,60); if the observed diffusion time (τobs) does not scale with the size of the excitation volume, then τobs is solely determined by the residence time tbound (≈1/koff) on the immobile structure (Figure 3A). If however τobs does scale with the size of the excitation volume, the binding process is much faster than the diffusion time scale (Figure 3B). Put differently, the relaxation time of the binding process [−1/(k⁺koff)] will be smaller than τobs if τobs varies with the focal spot size. We first performed detailed in vitro control experiments to characterize this set-up (Figure 3C). In aqueous solution, both the apparent particle number (Supplementary Figure S4A) and the average diffusion time (Supplementary Figure S4B) of the standard probe rhodamine 6G increased when tuning the laser beam to a smaller diameter (and hence the focal spot to a larger diameter) (Figure 3D black). In a next step, we used eGFP as a probe. Both in phosphate buffered saline (PBS) (Figure 3D, magenta) and in a buffer with an index of refraction matching the intracellular environment (PBS, 23.5% w/w sucrose, n = 1.37) (Figure 3D, green) increasing the laser focus again shifted the experimental ACF to a slower time scale, as expected. The TFFCS set-up thus performs well for determining the dependence of the ACF on the size of the focus. We next performed similar measurements in living HeLa cells (Figure 3E). The diffusion time of eGFP was clearly dependent on the size of the focal spot.

In summary, we combined sFRAP, CP and FCS to study the mobility of eGFP-LEDGF/p75 in the nucleus of living HeLa cells. Overall, LEDGF/p75 does not interact strongly with chromatin, as implied from the absence of a permanent immobile fraction in sFRAP. Rather, we observed different dynamic states: (i) a slow state with D = 0.4–0.5 μm²/s, observed with sFRAP and FCS, (ii) a presumed bound state with ~60 nM binding sites and a dissociation constant Kdiss ~100 nM, suggested from sFRAP and CP measurements and (iii) a fast state, observed only with FCS.
Figure 3. TFFCS shows fast chromatin binding of LEDGF/p75. (A) The observed slow diffusion time ($\tau_{\text{obs}} = 25.9$ ms, $D = 0.5 \mu m^2/s$) is the binding time of LEDGF/p75 with chromatin, $t_{\text{bound}} = 1/k_{\text{off}}$. In this case $\tau_{\text{obs}}$ does not scale with the laser focus diameter, indicated in blue. (B) LEDGF/p75 continuously associates with and dissociates from the chromatin while diffusing through the measurement spot. The $\tau_{\text{obs}}$ scales with the laser focus diameter. (C) Schematic representation of the home built TFFCS set-up. The beam expander and diaphragm are used to vary the diameter of the collimated laser beam. ND, neutral density attenuation filter; APD, avalanche photodiode. The diameter of the diaphragm is controlled by measuring the power of the excitation light passing through it. Higher power through the diaphragm means a broader beam will enter the objective back aperture. The objective focuses a broader beam in a smaller focal point, since the beam diameter determines the effective numerical aperture of the objective ($NA = n \times \sin \alpha$ with $n$ the refractive index and $\alpha$ the angular aperture). For further details, see the ‘Materials and Methods’ section. (D) In vitro TFFCS measurements of rhodamine 6G in water (black), eGFP in PBS (magenta) and PBS/sucrose with a viscosity and refractive index similar to that of the intracellular environment ($n = 1.37$) (green). $\omega_0$: radial radius of the excitation spot. (E) Intracellular TFFCS measurements of eGFP (black), eGFP-LEDGF/p75 (magenta) and eGFP-LEDGF/p75 K56D-R74D (green). Error bars = SD. (F–H) Plot of the diffusion time from the experimental ACF versus the squared radial radius of the confocal excitation spot for (F) eGFP, (G) the fast and (H) the slow component of eGFP-LEDGF/p75.
LEDGF/p75 shows normal diffusion dependent dynamics. (A) Illustration of half-nucleus FRAP. Half of the nucleus is photobleached and recovery perpendicular to the bleach border is monitored versus time. (B) Nuclear intensity profile versus time along the arrow designated ‘position’ in (A) Solid lines = Equation 10. (C) Mean squared displacement – time plot. Error bar = SD. Solid line = Equation 11.
eGFP-LEDGF/p75 K56D-R74D still varied with the focus size, implying that this protein also moves by diffusion (Figure 3E). In summary, by mutating a single amino acid residue in LEDGF/p75-PWWP, R74D, chromatin binding of the protein was severely impaired, as evidenced by an up to 8-fold increase in dynamics with respect to wild-type LEDGF/p75. The PWWP domain of LEDGF/p75 is thus important but not sufficient for in vivo chromatin binding.

LEDGF/p75 locks the IN-LEDGF/p75 complex on chromatin through its PWWP domain

LEDGF/p75 has been shown to tether IN to chromatin by virtue of a direct protein–protein interaction (15,22,61–64). We performed CP measurements of eGFP-tagged IN in HeLa cells to show tethering in vivo: we observed variable photobleaching of eGFP-IN from cell to cell in wild-type HeLa cells (Figure 6A), but no significant photobleaching of eGFP-IN was observed in HeLa-p75KD cells (Figure 6B). When HeLa-p75KD cells were back-complemented with 500 nM of RNAi-resistant mRFP-LEDGF/p75, a large fraction of eGFP-IN was again photobleached (Figure 6C). The dynamics of IN are thus strongly influenced by LEDGF/p75 mediated chromatin tethering. We next quantified the effect of IN on in vivo chromatin binding kinetics of LEDGF/p75. Upon co-expression of mRFP-labeled IN the overall dynamics of LEDGF/p75 shifted to much slower time scale (Figure 6D). When fitting the sFRAP curve with a normal diffusion model (Supplementary Equation S1), a good fit was obtained (Figure 6D and Supplementary Figure S5A) and \( D = 0.0055 \pm 0.001 \text{m}^2/\text{s} \) was calculated, about 75 times lower than in cells without co-expression of mRFP-IN (Tables 1 and 3). A reaction-dominant model (Supplementary Equation S3) did not give a good fit (Supplementary Figure S5B) and a full diffusion-reaction model (Supplementary Equation S4) did not result in a better fit than a normal diffusion model (Supplementary Figure S5C). With hnFRAP a similar decrease in dynamics was observed (\( D = 0.021 \pm 0.004 \text{m}^2/\text{s} \)) (Figure 6E and F).

Next, we verified with sFRAP whether the mutations in the PWWP domain of LEDGF/p75 affected chromatin tethering of IN. Co-expression of mRFP-IN decreased the dynamics of the K56D-R74D mutant about six times, as calculated from the difference in D (Tables 2 and 3 and Figure 6G), which is much less than wild-type eGFP-LEDGF/p75 (note the difference in time scale in Figure 6D and G). To verify that the absence of chromatin tethering was not due to a loss of the IN-LEDGF/p75 interaction, we used cellular FCCS. A high relative cross-correlation amplitude, comparable with what we published before for IN-LEDGF/p75 K150A (61,62), was observed (Figure 6H), indicating that the interaction is not affected by the mutations. In conclusion, chromatin tethering of IN by LEDGF/p75 can be observed with FRAP. We showed for the first time in vivo that IN-LEDGF/p75 complexes have a 75-fold lower mobility as compared with LEDGF/p75. Since IN alone did not appear to
interact strongly with chromatin. LEDGF/p75 thus tethers IN strongly to chromatin. Rational mutation of the PWWP domain resulted in a strongly decreased ability to tether IN, indicating that the PWWP domain in LEDGF/p75 constitutes a ‘protein lock’ that strongly tethers IN to chromatin.

**DISCUSSION**

**Fluorescent LEDGF/p75 displays wild-type properties**

Our experimental approach combines complementary fluorescence techniques to study fluorescently labeled LEDGF/p75 in living cells. It had been shown that the interactions with cellular partners through the IBD are not hampered by the presence of the eGFP label (28–30,63). Unlike a C-terminal eGFP (data not shown), an N-terminal fusion displayed a proper intracellular distribution. In addition, the overall chromatin binding characteristics of eGFP-LEDGF/p75 are not altered (Figure 1E); since LEDGF/p75 lost chromatin binding by mutating the PWWP domain, we are confident that chromatin binding of eGFP-LEDGF/p75 is conserved. Results are in agreement with Turlure et al. (16), who showed that eGFP-PWWP still functionally interacts with chromatin. All experiments in cells were performed at relevant concentrations of fluorescent LEDGF/p75. On average, the concentration of transiently expressed eGFP-LEDGF/p75 was similar to that of endogenous LEDGF/p75 (Figure 1D) and measurements were performed in a 1 nM–2.5 μM concentration range to study potential concentration dependent effects (Figure 1D). Furthermore, the ACF of eGFP-LEDGF/p75 was concentration independent (data not shown) and our investigation of chromatin tethering of IN (Figure 6A–C) led us to assume that the intracellular concentration of LEDGF/p75 lies in submicromolar range. In line with this, LEDGF/p75 has been shown to be a much more abundant cellular protein than its alternative splice variant p52 (22,65).

**Dynamic in vivo chromatin interactions of LEDGF/p75**

DNA binding and chromatin fractionation assays have been used in the past to study interactions of LEDGF/p75 with chromatin but these methods only provide a static and average view on interactions with DNA/chromatin (15,16,21). Also, irreconcilable notions of in vivo chromatin binding of LEDGF/p75 exist. On the one hand, LEDGF/p75 has been shown to be a ‘site-specific’ transcriptional co-activator, interacting with promoter elements of stress-related genes. On the other hand, LEDGF/p75 has been shown to be the factor that targets lentiviral integration to ‘random’ genomic regions of active transcription (21,24). We used sFRAP, CP, FCS, TFFCS and hnFRAP to monitor in vivo chromatin interactions of LEDGF/p75 for the first time. We showed that LEDGF/p75 does not permanently take part in immobile protein/chromatin complexes (Figure 2A).

The slow state was observed both with all techniques and represents low-affinity, likely non-specific chromatin scanning/hopping, a common phenomenon of transcription factors (66). By this mechanism, LEDGF/p75 likely targets lentiviral integration to random (actively transcribed) regions (24–26). On the other hand, the concentration dependent CP measurements strongly suggest the existence of low-concentration higher-affinity chromatin-bound states (Figure 2A–C and Table 1), which might result from association with promoters of stress-responsive genes (21,67) or more generally, as has recently been shown, from the association downstream of transcription start sites (TSS) of active transcription units (68). The concentration of binding sites we calculated (61 nM), indeed lies on the same order-of-magnitude as the concentration of TSS in an eukaryotic cell (~5–100 nM, see ‘Materials and Methods’ section).

To our surprise, we only observed the fast state (Figure 2D and Table 1) with FCS. This state did not appear to represent translational motion of the protein (Figure 3G). The time scale and fraction of the fast process did not appear to be concentration or cell dependent (data not shown), ruling out free/bound protein fractions or a post-translationally modified fraction. This state might represent any process that changes the emissive state of the eGFP fluorophore, such as protonation (69), quenching or possibly rotational diffusion of chromatin bound eGFP-LEDGF/p75. In that respect, it has been shown that mRFPs can reside in dark states even in the milliseconds time scale under 1P excitation (70).

### Table 2. Dynamics of LEDGF/p75 PWWP mutants measured with sFRAP and FCS

| Mutant          | sFRAP | FCS | sFRAP | FCS | sFRAP | FCS |
|-----------------|-------|-----|-------|-----|-------|-----|
| LEDGF/p75       |       |     |       |     |       |     |
| eGFP-LEDGF/p75  |       |     |       |     |       |     |
| K56D            | F_0m  | 0.1 ± 0.1 | F_0m  | 0.1 ± 0.1 | F_0m  | 0.1 ± 0.1 |
| D_0m (μm^2/s)   | 16.4 ± 3.5 | 13.3 ± 2.4 | 16.8 ± 5.1 |
| F_0f (%)        | 44.5 ± 9.3 | 58.5 ± 8.6 | 62.8 ± 13.8 |
| D_0f (μm^2/s)   | 1.97 ± 0.09 | 1.0 ± 0.3 | 1.5 ± 0.8 |
| α               | 0.69 ± 0.05 | 0.78 ± 0.04 | 0.80 ± 0.05 |

Values represent averages of at least 10 (sFRAP) or 20 (FCS) measurements and are ±SD. Parameters D_0m, F_0f and D_0f are obtained from a two-component normal diffusion fit. Parameter α is obtained from a one-component anomalous fit.

*Fit not satisfying. For more details on the parameters, see Table 1.
Quantitative information on in vivo chromatin hopping with TFFCS

We showed with hnFRAP that diffusion of eGFP-LEDGF/p75 in the slow (seconds) time scale seems normal in absence (Figure 4C) or presence (Figure 6F) of HIV-1 IN. By adapting a standard FCS microscope, we could also clearly distinguish the contribution of diffusion to the observed chromatin binding dynamics of LEDGF/p75 and showed that even in the fast (millisecond) time scale diffusion still seems normal (Figure 3E and H). This implies that association with and dissociation from chromatin occurs faster than the transit time of the protein through the laser focus, as illustrated in Figure 3B. As such, the strength by which LEDGF/p75 is slowed down could be used to quantify the low-affinity chromatin binding of LEDGF/p75. Based on $D_{\text{observed}} = 0.5 \mu m^2/s$.
and the theoretical $D_{\text{free}} = 22 \mu m^2/s$, we calculated with Equation 8 that LEDGF/p75 remains 98% of the time bound to chromatin and moves during the remaining 2% of the time to a new binding site. The average $t_{\text{free}}$ and $t_{\text{bound}}$ should thus be considerably shorter than the observed diffusion time (25.9 ms) and $k^*_{\text{on}} \sim 43$ times larger than $k_{\text{off}}$ (Equation 7). In other words, $k_{\text{off}}$ should be bigger than $38.6/s$ and $k^*_{\text{on}} > 1.7 \times 10^3/s$. This kinetic ‘effective diffusion’ regime of non-specific chromatin binding has been simulated before (36) and implies that transcription factor chromatin binding might be even more dynamic than currently appreciated (33,36,42,59).

Importantly, TFFCS as such would not be able to discern between slow movement of a (big) protein complex and frequent chromatin interactions of a (small) protein. Therefore, one has to verify that chromatin interactions are indeed contributing to the observed dynamics, which we did, both in vitro (Figure 1E) and in cells (Supplementary Figure S3). Furthermore, by rational mutation of the PWWP domain, a known chromatin interaction motif, we did prove that the PWWP contributes to the observed affinity of LEDGF/p75 for chromatin in vivo. Additionally, we could show cargo-mediated chromatin tethering by LEDGF/p75, which has been shown many times to be an important function of LEDGF/p75 for HIV replication. Finally, it must be noted that the ‘chromatin hopping’ we describe for LEDGF/p75 could also represent chromatin hopping of a larger complex containing LEDGF/p75 and other nuclear components and this complex would experience a larger effective viscosity in the chromatin matrix. Further research will provide more insight in this. In line with our kinetic regime of hopping, fast chromatin binding kinetics have recently also been experimentally observed for the *Drosophila* Hox gene Sex combs reduced transcription factor with DNA in vivo (60) and DNA-bound and slow apparent diffusion two-state chromatin binding has recently also been observed for Heterochromatin protein 1 (43), a protein we also studied as a control for chromatin binding (Supplementary Figure S3).

### Extremely slow, yet diffusion controlled dynamics of LEDGF/p75-IN

| eGFP-LEDGF/ p75 + mRFP-IN | eGFP-LEDGF/ p75 K56D-R74D+ mRFP-IN |
|-----------------------------|----------------------------------|
| sFRAP                      |      hnFRAP                      |
| $F_{\text{on}}$ (%)        | ~4.5 ± 2.0                       |
| $D$ (µm²/s)                | 0.006 ± 0.001 0.021 ± 0.004 0.54 ± 0.02 |
| $F_{\text{off}}$ (%)       | 0.12 ± 0.02                      |

Values represent averages of at least 10 (sFRAP) or 5 (hnFRAP) measurements and are ±SD. For more details on the parameters, see Table 1.

IN has been suggested to be present in a paracrystalline form in the viral particle (71). In the presence of its viral cargo, IN, LEDGF/p75 formed a 75-fold slower dynamical complex. Since interaction with IN likely does not decrease $k^*_{\text{on}}$ of LEDGF/p75, the observed dynamics should still be diffusion controlled, which is what we observed (Supplementary Figure S5 and Figure 6E and F). The effective diffusion law (Equation 7) allowed us to calculate that while for LEDGF/p75, $k^*_{\text{on}}/k_{\text{off}} \sim 43$, for IN-LEDGF/p75, $k^*_{\text{on}}/k_{\text{off}} \sim 2.7 \times 10^3$. The $k^*_{\text{on}}$ is thus at least as large as for LEDGF/p75 alone and the $k_{\text{off}}$ likely decreases considerably after interaction with IN. The IN is known to be present at least in a dimeric form at nM–µM concentrations (72) and already a IN dimer contains two binding sites for LEDGF/p75 (73). As we have shown, IN does not bind strongly to chromatin (Figure 6A–C). Probably thus, IN-LEDGF/p75 complexes contain at least two LEDGF/p75 units that can both tether IN cooperatively to chromatin, resulting in the apparent diffusion controlled increase in affinity. An allosteric effect of IN on the intrinsic affinity of LEDGF/p75 for chromatin is unlikely, since the IBD and chromatin binding domains of LEDGF/p75 are located in distinct structural parts of the protein. It might be argued that the IN-LEDGF/p75 complex senses, due to its oligomeric size, a much larger viscosity in cells than does LEDGF/p75 alone and this might explain the observed 75-fold decrease in dynamics. As we have shown in Figure 6G and H, PWWP-mutated LEDGF/p75 still has wild-type affinity for IN, but its dynamics are only slowed down 6-fold by IN co-expression. Therefore, we are confident that the observed dynamical shift of IN-LEDGF/p75 does represent an increased affinity for chromatin.

The linker histone H1, which we also used as a control for our measurements (Supplementary Figure S3), has recently also been studied quantitatively and similar slow overall dynamics were observed (74). Because dynamics of H1 were described best with a full diffusion/reaction model while a simple diffusion model sufficed for IN-LEDGF/p75, this suggests that while for H1 very slow dissociation from chromatin is the cause of the slow overall dynamics, very fast association with chromatin is rather the cause of the slow dynamics for IN-LEDGF/p75.

### The PWWP domain is the key to the chromatin lock

In recent years, LEDGF/p75 has become a protein of interest to understand various diseases such as leukemia, autoimmunity and AIDS (4,7,8,22). Nearly all normal and disease related functions of LEDGF/p75 involve tethering of interaction partners to chromatin through its conserved IBD. We investigated the role of the PWWP domain in the chromatin binding of LEDGF/p75 in vivo. The conserved PWWP domain of LEDGF/p75 contains a recently described general protein fold, present in proteins that carry chromatin-binding motifs and has been proposed to bind DNA in several independent structural biology studies (54,75). Other studies however suggested, albeit without experimental evidence so far, that the PWWP fold is implicated in non-DNA chromatin interactions.
(12,14). Although eGFP-PWWP did not show any DNA binding in vitro, it was shown to be localized on mitotic chromosomes, much alike full length LEDGF/p75 (16). Our rational mutagenesis of the PWWP domain of LEDGF/p75 resulted in a protein with up to 8-fold increase in overall mobility as compared with wild-type LEDGF/p75 (Figure 5D and F). The mutant protein exhibited diffusion-dependent anomalous diffusion (Figures 3E and 5F and Table 2), which could either result from residual interactions with the chromatin (76) or from interactions with other cellular cofactors, leading to obstructed diffusion (77). Nonetheless, we confirm that interactions of the PWWP domain with chromatin are necessary but not sufficient to convey wild-type affinity of LEDGF/p75 for chromatin.

Importantly, we are the first to show that upon mutation of the PWWP-domain, tight chromatin tethering of IN disappears, while the tight interaction of LEDGF/ p75 with IN is conserved. This strongly argues for a sole role of LEDGF/p75, in particular the PWWP-domain, in associating an IBD-bound cargo to chromatin. The HDGF-PWWP has been shown to be able to dimerize (78) and this might be an important step in chromatin tethering. Currently, we are investigating whether cargo-induced locking is a general phenomenon for tethering cargos to chromatin. Being the key for locking cargos to chromatin, the PWWP-domain might be a promising novel target for blocking LEDGF/p75 malfunctioning in AIDS but also oncogenesis.

SUPPLEMENTARY DATA
Supplementary Data are available at NAR Online.

ACKNOWLEDGEMENTS
The authors acknowledge Davide Mazza (Laboratory for Receptor Biology and Gene Expression, National Cancer Institute, Bethesda, MD) for providing the Matlab code for analyzing the FCS data with the binding models, Antoine Delon (Université Joseph Fourier, France) for helpful discussions on CP and Werner Verbakel (Laboratory for Biomolecular Dynamics) for writing the ImageJ script for hnFRAP analysis.

FUNDING
Funding for open access charge: University of Leuven (GOA/2006/02, IOF/KP/07/008); the Institute for the Promotion of Innovation through Science and Technology in Flanders (CellCoVir SBO grant 60813); the Research Foundation Flanders (FWO) (grant G.0530.08); the European Commission project Targeting HIV Integration Co-factors (grant HEALTH-F3-2008-201032); FWO postdoctoral fellowship (to J.Hendrix); Mathilde-Krim postdoctoral fellowship (The Foundation for AIDS Research) (to J.D.R.).

Conflict of interest statement. None declared.

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