Oxidation-induced Structural Changes of Ceruloplasmin Foster NGR Motif Deamidation That Promotes Integrin Binding and Signaling

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Significance: Asparagine deamidation at Asn-Gly-Arg (NGR) sites leads to the isoAsp-Gly-Arg (isoDGR) integrin-binding motif formation.

Results: Ceruloplasmin (Cp), which contains two NGR sites and is oxidized in cerebrospinal fluid (CSF) in neurodegenerative diseases, can undergo oxidation-induced structural changes fostering NGR deamidation with gain of integrin binding and signaling properties, in vitro and ex vivo in pathological CSF.

Conclusion: Cp NGR motifs can deamidate acquiring integrin-binding functions.

Background: Asparagine deamidation at Asn-Gly-Arg (NGR) sites leads to the isoAsp-Gly-Arg (isoDGR) integrin-binding motif formation.

Asparagine deamidation occurs spontaneously in proteins during aging; deamidation of Asn-Gly-Arg (NGR) sites can lead to the formation of isoAsp-Gly-Arg (isoDGR), a motif that can recognize the RGD-binding site of integrins. Ceruloplasmin (Cp), a ferroxidase present in the cerebrospinal fluid (CSF), contains two NGR sites in its sequence: one exposed on the protein surface (568NGR) and the other buried in the tertiary structure (962NGR). Considering that Cp can undergo oxidative modifications in the CSF of neurodegenerative diseases, we investigated the effect of oxidation on the deamidation of both NGR motifs and, consequently, on the acquisition of integrin binding properties. We observed that the exposed 568NGR site can deamidate under conditions mimicking accelerated Asn aging. In contrast, the hidden 962NGR site can deamidate exclusively when aging occurs under oxidative conditions, suggesting that oxidation-induced structural changes foster deamidation at this site. NGR deamidation in Cp was associated with gain of integrin-binding function, intracellular signaling, and cell pro-adhesive activity. Finally, Cp aging in the CSF from Alzheimer disease patients, but not in control CSF, causes Cp deamidation with gain of integrin-binding function, suggesting that this transition might also occur in pathological conditions. In conclusion, both Cp NGR sites can deamidate during aging under oxidative conditions, likely as a consequence of oxidative-induced structural changes, thereby promoting a gain of function in integrin binding, signaling, and cell adhesion.
NGR motif, and ~0.5% contain more than one NGR motif, suggesting a regulated functional role for this motif. Nevertheless, depending on the protein sequences and structures and on environmental conditions, only certain NGR sites with suitable features are likely to undergo deamidation.

The copper protein ceruloplasmin (Cp), a ferroxidase enzyme present in the cerebrospinal fluid (CSF) (15, 16), contains two NGR sites (568NGR and 962NGR). We previously reported that Cp undergoes oxidative modifications in the CSF of Parkinson disease and AD patients (17), as a consequence of changes in the environmental redox status of pathological CSF (18, 19). Cp oxidation causes a decrease in its ferroxidase activity that may have pathological implications (17). Because oxidative conditions might cause structural changes that accelerate Asn deamidation, we investigated whether the Cp NGR motifs can deamidate during aging under oxidative conditions and whether, as a consequence, Cp acquires integrin binding properties mediated by isoDGR.

Here, we show that although both Cp NGR sites can deamidate, the 962NGR motif undergoes deamidation only when Asn aging occurs under oxidative conditions, in which structural changes are induced. The NGR to isoDGR transition in Cp induces gain of integrin binding function, integrin-mediated intracellular signals, and cell pro-adhesive activity. Finally, we show that Cp deamidation is faster in the CSF from AD patients compared with the one from healthy subjects, suggesting that this Cp modification, and the consequent gain of function, might occur in vivo in pathological conditions.

**EXPERIMENTAL PROCEDURES**

**Patients**

Samples were obtained from the Institute of Experimental Neurology INSPE-Biobank (San Raffaele Scientific Institute, Milan, Italy). After approval from the hospital’s ethical committee and informed consent from patients, CSF samples (0.8–1 ml) were collected by lumbar puncture. The analyzed groups were: Alzheimer disease patients (n = 16) (Table 1). All patients were at first diagnosis and drug-free. Current criteria for the diagnosis of AD (20) were used for patients admission into the study. Exclusion criteria were: HIV or Hepatitis C virus seropositivity; the appearance of other neurodegenerative diseases or previous cerebral ischemic events; and severe metabolic disorders. Control CSF was from patients who underwent lumbar puncture on account of a suspected neurological disease and who proved to be normal and free from pathological alterations after complete CSF analysis and thorough clinico-neuroimaging assessment. Samples were centrifuged (800 × g, 10 min at 4 °C) to eliminate cells, then were either immediately processed, or stored at −80 °C in an N2-supplemented atmosphere to avoid oxidation.

**Cell Cultures and Reagents**

Human glioblastoma T98G, keratinocyte HaCaT, and endothelial EA.hy926 cell lines (ATCC) were cultured (37 °C in a 5% CO2 atmosphere) in DMEM (ATCC) supplemented with L-glutamine, and 10% FBS. Human integrins α5β1, αvβ3, and αvβ5 (Immunological Sciences), recombinant αvβ6 and αvβ8 (R&D System), and human plasma purified Cp (Alexis Biochemicals) were used.

**Oxidation and Asparagine Accelerated Aging Treatments**

Oxidation and accelerated Asn aging treatments were performed by incubating purified Cp at 37 °C in various buffers as described below. Oxidized Cp was prepared as follows: purified Cp was diluted at 1 mg/ml in PBS buffer containing 10 mM H2O2 solution and incubated for 16 h at 37 °C. Deamidated Cp was prepared as follows: purified Cp was diluted 1 mg/ml in 100 mM ammonium bicarbonate (AmBic) buffer, pH 8.5, incubated for 16 h at 37 °C, and stored at −20 °C until analysis. This condition is known to favor Asn deamidation at the NGR site (10). Before use in the assays, Cp was dialyzed against PBS using Slide-A-Lyzer dialysis cassette 10,000 molecular weight cutoff (Pierce). Oxidized and deamidated Cp was obtained by combining the above treatments. These products are hereinafter

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**TABLE 1**

Demographic and clinical features of patients and controls

|   | Sex | Age | [C] | Sex | Age | [C] | MMSE |
|---|-----|-----|-----|-----|-----|-----|------|
| CN | M   | 70  | 0.37| F   | 56  | 0.41| 22   |
|   | M   | 68  | 0.32| M   | 75  | 0.38| 18   |
|   | M   | 71  | 0.38| M   | 71  | 0.39| 21   |
|   | M   | 62  | 0.16| M   | 73  | 0.38| 18   |
|   | M   | 63  | 0.85| M   | 72  | 0.33| 7    |
|   | M   | 54  | 0.25| F   | 72  | 0.41| 18   |
|   | F   | 72  | 0.40| M   | 69  | 0.29| 18   |
|   | F   | 77  | 0.50| M   | 64  | 0.33| 18   |
|   | M   | 61  | 0.34| F   | 73  | 0.45| 13   |
|   | M   | 70  | 0.28| M   | 79  | 0.61| 24   |
|   | F   | 72  | 0.41| F   | 78  | 0.33| 21   |
|   | F   | 77  | 0.50| F   | 77  | 0.33| 16   |
|   | M   | 80  | 0.40| M   | 70  | 0.24| 25   |
|   | F   | 84  | 0.23| M   | 77  | 0.41| 27   |
|   | M   | 81  | 0.29| F   | 79  | 0.53| 29   |
|   | M   | 81  | 0.39| M   | 86  | 0.49| 19   |

*These values are the means ± S.D.
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referred to as “oxidized Cp” (Cp-ox), “deamidated Cp” (Cp-AmBic), and “oxidized/deamidated Cp” (Cp-ox/AmBic), respectively. Aging of Cp in CSF was performed by adding purified Cp (final concentration, 20 μg/ml) to CSF either from healthy subjects or from AD patients. CSF samples were then left to incubate for different time (0, 3, 6, 9, and 12 days) at 37 °C under nitrogen conditioned atmosphere, to avoid the exposure to atmospheric oxidative environment.

Binding of Cp to Integrins and Competition with isoDGR Peptide

α5β1, αvβ3, αvβ5, αvβ6, and αvβ8 integrins (1 μg/ml in PBS with Ca2+/Mg2+; DPBS, Cambrex) were added to 96-well polyvinyl chloride plates (BD Biosciences) and incubated 12 h at 4 °C. Subsequent steps were carried out at 20 °C. After blocking (3% BSA-PBS), the plates were filled with: Cp solutions (2–20 μg/ml in binding buffer; 25 mM Tris-HCl, pH 7.4, 150 mM NaCl, 1 mM MgCl2, 1 mM MnCl2, 0.05% Tween, 1% BSA) or CSF samples from healthy subjects or AD patients supplemented with Cp (prepared as described above, diluted in binding buffer) and incubated for 2 h. Binding was detected using a polyclonal anti-human Cp Ab (Abcam) followed by a secondary HRP conjugate Ab (Abcam) and by o-phenylenediamine chromogenic substrate. Competitive binding assays were performed by mixing 10 μg/ml of either the acetyl-CisoDGRCGVSSRTPSDKY peptide (isoDGR peptide) or the control peptide CARACGVRSSRTPSDKY (ARA peptide) (12) with Cp-ox/AmBic solutions (20 μg/ml) and mixing isoDGR peptide (30 μg/ml) with CSF samples spiked with Cp (20 μg/ml). The mixtures were added to microtiter plates coated with integrin, and the binding assay was carried out as described above. Cp bound to the αvβ6-coated plates was analyzed as follow; after the binding, the plates were washed and incubated with Laemmli buffer (5 min at 100 °C) to promote protein detachment, the solution was then collected and subjected to SDS-PAGE and Western blot analyses as described (17) using a polyclonal anti-human Cp Ab (Abcam). Images were acquired using a laser densitometer (Molecular Dynamics).

Cell Adhesion Assay and Protein l-Isoaspartyl Methyltransferase (PIMT) Treatment

96-well polyvinyl chloride microtiter plates were coated with either untreated or treated Cp or with Cp that was immunoprecipitated (see below) after aging in healthy or AD CSF (1–20 μg/ml in 50 mM Na3PO4, pH 7.3, 150 mM NaCl, 16 h at 4 °C). After washing and blocking with 3% BSA-DPBS, the plates were seeded with HaCaT, EA.hy926, and T98G cells diluted in 0.1% BSA-DMEM (40,000 cells/well) and left to adhere for 3 h at 37 °C. Adherent cells were fixed and stained with crystal violet as described (14), and adhesion was evaluated as absorbance at 570 nm. For each treatment condition, cell spreading was evaluated comparing cell area in different microscopy fields. To measure cell area, pixels were automatically detected by image software analysis (Progenesis PG240; Nonlinear Dynamics). For PIMT treatment, plates coated as described above, were washed and filled with 45 μl of 0.02 mM S-adenosyl-l-methionine in 50 mM Na3PO4, pH 6.8, and 5 μl of PIMT solution (from IsoQuant isoaaspartate detection kit; Promega) and incubated at 37 °C for 16 h. Then plates were washed, and cell adhesion assay was performed as described above.

Cp Immunoprecipitation from CSF

After incubation in healthy or AD CSF, spiked purified Cp was immunoprecipitated using protein G-agarose beads (Invitrogen) coated with an anti-Cp antibody (Abcam ab8813). The antibodies were cross-linked to the beads with 20 mM dimethyl pimelimidate (Sigma) and added to the CSF supplemented with Cp; the solution was then incubated for 24 h at 4 °C under gentle stirring. Beads were washed, and Cp was eluted with 0.1 M glycine solution, pH 2.5; samples were immediately diaphragmed with PBS using an Amicon filter device.

Bathophenanthroline Assay

Ferrooxidase activity of Cp after aging in CSF was evaluated on immunoprecipitated Cp by bathophenanthroline (Btp) assay as in ref. 17. Briefly, immunoprecipitated Cp (1.25 μg) was incubated with 80 mM FeSO4 (ferrous form) and analyzed after 1 h with a solution of 1 mM Btp in acetate buffer, pH 6.2. A decrease in Btp-Fe2+ complex absorbance at 535 nm derives from ferrous iron oxidation into ferric form (Fe3+).

Mass Spectrometry Analysis

Samples were dialyzed against digestion buffer and incubated 2 h at 20 °C with trypsin (10 ng/μl; Roche Applied Science). Digested samples were desalted (Stage tips C18; Thermoscientific) and injected in a capillary chromatographic system (EasyLC; Proxeon Biosystem). Peptide separations occurred on a 25-cm reverse phase silica capillary column, packed with 3-μm ReproSil 100 Å C18 AQ. A gradient of acetonitrile eluents was used to achieve separation (flow rate, 0.15 μl/min). MS analysis was performed by nanoLC-MS/MS using an LTQ-Orbitrap (ThermoScientific) equipped with a nanoelectrospray ion source (Proxeon Biosystems). Full scan spectra were acquired with the lock mass option, resolution set to 60,000, and mass range from m/z 350 to 1700 Da. The six most intense doubly and triply charged ions were selected and fragmented in the ion trap. All MS/MS samples were analyzed using Mascot (v.2.2.07; Matrix Science) and X!Tandem (within Scaffold software, v.2.06_00, 2007; Proteome Software Inc.) search engines to search the UniProt_Human Complete Proteome_c pt_hum_2012_07. Searches were performed with three missed cleavages allowed, N terminus acetylation, methionine oxidation, and deamination of asparagine/glutamine as variable modifications. Mass tolerance was set to 5 ppm and 0.6 Da for precursor and fragment ions, respectively. Scaffold was used to validate MS/MS-based peptide and protein identifications. Protein thresholds were set to 99.0% and two peptides minimum, whereas peptide thresholds were set to 95% minimum.

Circular Dichroism and Melting Curves

CD spectra were acquired for Cp and Cp after oxidative and aging treatments (at 1 μM in 50 mM PBS, pH 7) on a Jasco J-815 CD spectrometer at 20 °C. Each spectrum was averaged using four accumulations collected in 0.1-nm intervals with an average time of 0.5 s. The protein spectra were corrected by sub-
tracting the corresponding buffer spectra and then smoothed. The observed ellipticity (mdeg) was converted into molar residue ellipticity \( [\theta] \) \( \text{deg cm}^2 \text{dmol}^{-1} \). Melting curves were calculated recording CD spectra in continuous from 20 to 96 °C \( (1 \text{ °C intervals}) \) in a wavelength range from 190 to 260 nm to achieve the better resolution. Melting temperature \( (T_m) \) was calculated by nonlinear fitting Boltzmann sigmoidal with Prism V4.03 software (GraphPad Inc.).

**Molecular Dynamics Simulations and Docking Calculations**

**Homology Modeling**—I-TASSER server was used to predict the secondary and tertiary structure of \( \alpha \bar{v} \beta 6 \) integrin (21).

**Molecular Dynamics (MD) Simulations**—MD simulations were performed on Cp wild type (P00450; Protein Data Bank code 2J5W), using the GROMACS 4.5.4 package (22) with the optimized parameters for liquid simulation force field. The \( 568 \text{isoDGR-Cp} \) and \( \alpha \bar{v} \beta 6 \) systems have been simulated in the same manner. Three independent 50-ns-long MD simulations (150-ns production run) were performed for \( \alpha \bar{v} \beta 6 \) to get a set of structures to be used in docking calculations. All the analysis were performed using the GROMACS utilities on the last 40 ns of each simulation concatenated in a single trajectory \( (120 \text{ ns total}) \). In particular, cluster analysis were performed using the Gromos algorithm.

**Docking Calculations**—Docking calculations of \( 568 \text{isoDGR-Cp} \) on the globular head of the extracellular part of \( \alpha \bar{v} \beta 6 \) have been performed using the docking program HADDOCK2.0 (23) with the optimized parameters for liquid simulation force field. We have docked a bundle of 27 \( 568 \text{isoDGR-Cp} \) structures onto a bundle of 10 \( \alpha \bar{v} \beta 6 \) structures, which correspond to the centers of the clusters obtained from MD simulations. The protocol follows a three-stage docking procedure: (a) randomization of orientations and rigid body minimization, (b) simulated annealing in torsion angle space, and (c) refinement in Cartesian space with explicit water. Ambigious interaction restraints \( (\alpha \bar{v} \beta 6 \) residues 150, 218, 506, 547, 548, 601, and 815; \( 568 \text{isoDGR-Cp} \) residues 568–570) were derived from the known interactions of the isoDGR motif of the cyclic peptide with \( \alpha \bar{v} \beta 6 \) (11). The best 200 solutions in terms of intermolecular energies were selected for a semiflexible simulated annealing in which the side chains of \( \alpha \bar{v} \beta 6 \) and of the \( 568 \text{isoDGR-Cp} \) located at the binding interface were allowed to move in a semi-rigid body docking protocol to search for conformational rearrangements. The models were then subjected to a water refinement step. The single best docked solutions were analyzed according to hydrogen bonds, salt bridge contact, and buried surface accessibility.

**Statistical Analysis**

Categorical data were analyzed by using Fisher’s exact test and two-tailed \( p \) value. Continuous data were evaluated by unpaired Student’s \( t \) test, if the data passed the normality test for Gaussian distribution as assessed by the Kolmogorov-Smirnov test or were evaluated by Mann Whitney test; a two-tailed \( p \) value was used for the comparison of two means and standard error. In all analyses, \( p < 0.05 \) was considered to be statistically significant. The analysis was performed with Prism V4.03 software (GraphPad Inc.).

**RESULTS**

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**Accelerated Aging of Cp under Oxidative Conditions Causes Deamidation of NGR Sites**—Sequence analysis of purified Cp by tandem MS showed that both \( 568 \text{NGR} \) and \( 962 \text{NGR} \) motifs were not deamidated (Fig. 1A and B, untreated) and that protein oxidation by incubation in 10 mM \( \text{H}_2\text{O}_2 \) at 37 °C cannot convert NGR to DGR/isoDGR (Fig. 1A and B, ox). In contrast, protein incubation in 100 mM AmBic or AmBic plus 10 mM \( \text{H}_2\text{O}_2 \), i.e., solutions that respectively mimic accelerated asparagine aging conditions (AmBic) (10) and accelerated asparagine aging under oxidative conditions (ox/AmBic), the \( 568 \text{NGR} \) motif was partially deamidated (Fig. 1A). Interestingly, the \( 962 \text{NGR} \) motif was partially deamidated exclusively following Cp aging under oxidative conditions (Fig. 1B ox/AmBic). The presence of Asp in place of Asn in the AmBic-treated Cp indicates that a deamidation reaction occurred at the two NGR sites of Cp, although MS did not ascertain whether DGR or isoDGR isoform was present.

A possible explanation for the different behavior of the \( 568 \text{NGR} \) and \( 962 \text{NGR} \) motifs may arise from analysis of the crystallographic Cp structure (Protein Data Bank code 2J5W) (25), which reveals that the \( 568 \text{NGR} \) motif is exposed on the surface of the protein, whereas the \( 962 \text{NGR} \) motif is less exposed and therefore less accessible to the solvent (Fig. 2A and B). The two sequences display differing tertiary contexts (Fig. 2C and D) that might in turn influence deamidation rates differentially. We then attempted to clarify the structural differences between the two sequences by a computational approach. Molecular dynamics simulations on wild type Cp (Protein Data Bank code 2J5W) showed a relatively stable structure with large fluctua-
tions of loop regions (Fig. 2E). Notably, the root mean square fluctuations, as calculated throughout the simulation, suggest that the two NGR sequences are more rigid than the other residues (Fig. 2F). Importantly, the 568NGR sequence is steadily accessible to the solvent throughout the simulation, whereas 962NGR is buried inside the protein and is blocked in a stable conformation by polar interactions with neighboring amino acids (Fig. 2C and D). Accordingly, we hypothesized that oxidation might affect the Cp structure and promote the exposure of the 962NGR motif, which in turn adopts a favorable conformation for deamidation. Indeed, it has been previously inferred that Cp structure is affected by oxidation (17, 26–28).

Cp Oxidation Induces Secondary Structure Changes—CD spectra obtained for purified Cp indicated that the secondary structure of Cp, which is principally composed of \( \beta \)-strands (45.5%, as evaluated by Uniprot tool), was partially affected by Cp-AmBic treatment (Fig. 2G, black line versus blue line); notably, large spectra changes occurred upon Cp-ox and -ox/AmBic treatment, suggesting structural alterations (Fig. 2G, black line versus green and orange lines). Interestingly, the spectra profile...
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FIGURE 2. NGR motifs structural analysis, Cp molecular dynamics modeling, and circular dichroism analysis indicate that $^{568}$NGR is buried inside Cp structure and that secondary structure changes induced by oxidative conditions might allow its surface exposure. A and B, surface representation of Cp. In red are highlighted the $^{568}$NGR and $^{962}$NGR sequences, respectively, as calculated by the GROMACS g_sas tool on crystallographic Cp structure (Protein Data Bank code 2J5W). The higher solvent-accessible surface of $^{568}$NGR is represented by the larger size of the cartoon representation of $^{568}$NGR and $^{962}$NGR sequences in Cp. The NGR amino acids are shown as orange sticks. The hydrogen bonds are shown as red lines.

E, cartoon representation of $^{568}$NGR and $^{962}$NGR sequences in Cp. The NGR amino acids are shown as orange sticks. The hydrogen bonds are shown as red lines.

G, the molecular dynamics simulations (3 × 50 ns, 150-ns total simulation time). $^{568}$NGR and $^{962}$NGR are indicated with red circles. The plot shows large fluctuations localized in the loop regions, notably the two NGR sequences show a rigid behavior, with low root mean square fluctuations values along the whole simulation.

Each spectrum was averaged using four accumulations collected in 0.1-nm wavelength intervals with an average time of 0.5 s. The observed ellipticity (mdeg) was converted into molar residue ellipticity (θ) (deg cm$^2$ dmol$^{-1}$). For each treatment, the melting temperature ($T_m$) is also indicated.
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Cp to bind αvβ6 integrin. However, the experimental evidence reported in Fig. 2 suggests that the deamidation of 562-NGR in oxidative conditions causes a considerable structural rearrangement that cannot be predicted by computational methods. Thus, we decided to perform molecular dynamics simulation only on 568isoDGR-Cp protein to generate a set of reliable structures to be docked onto αvβ6.

Cluster analysis on the 568isoDGR-Cp molecular dynamics allowed extraction of a number of representative structures of the simulations. A bundle of 27 isoDGR structures, corresponding to the center of the clusters, were docked onto a bundle of 10 αvβ6 structures, which are the center of the clusters obtained from 10 ns of molecular dynamics of the free αvβ6. To restrict the number of possible docked solutions, we defined a set of restraints (Cp-isoAsp568, αvβ6-Asp150, Asp218, Ala143, Pro184, Phe185, Thr238, Cp-Gly569 : αvβ6-Asp150, Asp218, Ala143, Pro184, Phe185, Thr238, and Cp-Arg179 : αvβ6-Arg179, Asp181, Ala143, Pro184, Phe185, Thr238) corresponding to canonical RGD or isoDGR integrin-binding site (11) as suggested from competition experiments. The docking calculations led to different solutions (Fig. 4A) whereby the cluster 3, representing the best solution, showed that the 568isoDGR-Cp docked onto αvβ6 integrin recapitulates the classical RGD/isoDGR receptor interactions (11) and creates some additional interactions (αvβ6/Cp: Ser199/β6/Glu437, 1.7 ± 0.1 Å; Tyr178 αv/Arg570, 2.1 ± 0.1 Å; Asp146 β6/Lys378, 1.9 ± 0.3 Å; Thr238 β6/5681As, 1.9 ± 0.2 Å; Asp150 αv/Arg570, 1.8 ± 0.2 Å; Asp147 β6/Lys562, 1.9 ± 0.3 Å) (Fig. 4B).

FIGURE 3. Cp aged under oxidative conditions is able to bind integrins via isoDGR motifs. A, screening by ELISA of the binding to several integrins of Cp under resting conditions (Untreated), after Asn accelerated aging (AmBic), or after Asn accelerated aging under oxidative conditions (ox/AmBic). B, analysis of the dose-dependent binding to αvβ6 of untreated Cp or aged/oxidized Cp (ox/AmBic). C, competitive binding of aged/oxidized Cp (Cp-ox/AmBic) to αvβ6 with the isoDGR peptide and control ARA peptide. D, Western blot analysis of the Cp-ox/AmBic compared with the untreated Cp (Controls) and of the Cp-ox/AmBic eluted from the ELISA microplate either coated (+) or noncoated (−) with αvβ6. All the assays were performed in triplicate for three independent experiments (n = 3). Statistical significance reported as p values was evaluated by Student’s t test. ****, p < 0.0001; ***, p < 0.001; **, p < 0.01; *, p < 0.05.

FIGURE 4. Molecular docking model of isoDGR-Cp/αvβ6 integrin binding site. A, HADDOCK scores versus root mean square deviations from the lowest energy complex structure for docking run performed of Cp-αvβ6. Different clusters are shown (circles) represented in different colors. They were obtained by fitting the models on the backbone of the αvβ6-binding site and then calculating the root mean square deviation values from the lowest energy solutions over the heavy atoms of the Cp protein. Structures belong to the same cluster if they differ by less than 2 Å in the pairwise root mean square deviation matrix. The HADDOCK score corresponds to the weighted sum of different energy terms (van der Waals, electrostatic, and restraint energies). Cluster 3 is the best solution that is represented in B. HADDOCK model of isoDGR-Cp/αvβ6 integrin binding site. Surface representation of αvβ6 binding pocket (green) in complex with isoDGR-Cp (cyan cartoon). The side chains of the residues forming stable interactions (as reported in text) are shown in licorice. Orange spheres correspond to Ca2+ ions; magenta spheres correspond to Ca2+ cation; green spheres correspond to Na+ ions. Red dotted lines denote the hydrogen bonds of the Cp with αv and β6 subunits. In magenta licorice is shown the 562-NGR motif. The arginine of 568isoDGR predominantly interacts with Asp560 of αv domain, whereas the same arginine can transiently interact with Asp574. The isoaaspartyl residue of isoDGR interacts with Ca2+ MIDAS ion located in the β6 domain (see zoomed area). Other interactions between 568isoDGR-Cp and αvβ6 have been highlighted; in particular we noticed a strong and stable network of salt bridges between Lys562 and Lys569 of isoDGR-Cp and Asp146 and Asp147 of β6 domain of αvβ6 integrin (interaction reported in text).
Deamidated Cp Mediates Cellular Adhesion and Spreading—The binding to integrins observed in vitro might not reflect the real binding properties of deamidated Cp, given the possibility that adsorption of purified integrins on microtiter plates might alter their conformation. Furthermore, we have previously shown that the isoDGR site of fibronectin but not its corresponding NGR site can promote cell adhesion (10–12, 14). To assess whether Cp can generate an isoDGR motif able to recognize integrins in physiological conformations, we analyzed the effect of Cp aged under oxidative conditions on the adhesion of human epithelial (HaCaT), endothelial (E.A.hy926), and glioblastoma (T98G) cell lines. Preliminary flow cytometry analysis showed αvβ6 expression by HaCaT, αvβ3 and αvβ5 expression by EA.hy926, and αvβ8 expression by T98G (data not shown).

Plates coated with ox/Ambic-Cp increased in a dose-dependent manner the adhesion and spreading of HaCaT cells, whereas untreated Cp induced little cell adhesion (Fig. 5A and B). In HaCat cells adhering to ox/ambic-Cp coated plates, cell spreading showed a significant (p = 0.0087) area increase (~20%) as compared with cells adhering to Cp coated plates (Fig. 5C). Similar pro-adhesive effects were observed with glioblastoma (Fig. 5D and E) and endothelial cell lines (data not shown).

To assess whether cell adhesion was mediated by isoDGR, we incubated ox/Ambic-treated Cp with PIMT, an enzyme that converts L-isoAsp residues to L-Asp (10, 30). PIMT treatment almost completely inhibited the pro-adhesive activity of ox/Ambic-treated Cp and of isoDGR peptide, the latter serving as a control (Student’s t test, p < 0.0001) (Fig. 5D and E). These results suggest that isoAsp formation, presumably at NGR site(s) of Cp, is associated with a “gain of function” in cell adhesion assays.

Cp Aged under Oxidative Conditions Transduces Intracellular Signal through Integrin Engagement—Having observed that Cp after accelerated Asn aging under oxidative conditions was able to bind integrins expressed on the cell surface, we investigated whether this binding could also induce intracellular signaling.Signal transduction pathway analysis, as performed by reverse phase protein arrays on epithelial cells, indicated that cell incubation with ox/Ambic-treated Cp can induce coordinated phosphorylation events that recapitulate many of the steps of classical integrin-mediated signal transduction. Hierarchical clustering analysis of the comparison of treatment with either ox/Ambic-Cp or untreated Cp revealed that after 30 min: (a) there was an increase in the phosphorylation of the activation residues of several molecules, e.g., p-Tyr997FAK1, p-Thr514PKCγ, p-Ser217/221MEK1, p-Thr185/187/202/Y397ERK1/2, p-Ser241/242PDK1, p-Ser473Akt, and (b) the phosphorylation of inhibitory residues, e.g., p-Ser259/320GSK3β, which in turn maintain β-catenin activity by preventing its phosphorylation (Fig. 6A and B). Several other proteins were unaffected (such as Crkl, SAPK/JNK, c-Jun, and Raf1) or slightly inhibited (PTEN and PKCa) (Fig. 6A and B). These results suggest that early signals mediated by deamidated Cp mostly addressed gene activation regulation, cell cycle, and MAPK signaling pathway (Fig. 7). In contrast, late signals (i.e., immediately after 120 min of treatment) seemed to sustain actin cytoskeleton rearrangement rather than cell survival, proliferation, and MAPK pathway activation (Fig. 7). Src and Crkl were activated, and the phosphorylation of FAK1 was inhibited, concomitantly with the slight phosphorylation of PTEN and Raf1 inhibitory molecules, which block MEK1 and ERK1/2 activation (Fig. 6A and B; Fig. 7). Similarly, Akt inactivation results in β-catenin inhibition by Ser25/37/45/Thr34 phosphorylation, as mediated by GSK3β activation (Fig. 6A and B; Fig. 7).

Pathological CSF from Alzheimer Disease Fosters Integrin Binding of Spiked Purified Cp and Inhibits Its Ferrooxidase Activity—To investigate whether Cp deamidation might occur also in vivo, in particular in pro-oxidant pathological CSF environments, we added purified Cp to CSF from healthy subjects or CSF from AD patients and incubated the mixture for 0, 3, 6, 12, 36, and 60 hours. The binding to integrins observed in vitro might not reflect the real binding properties of deamidated Cp, given the possibility that adsorption of purified integrins on microtiter plates might alter their conformation. Furthermore, we have previously shown that the isoDGR site of fibronectin but not its corresponding NGR site can promote cell adhesion (10–12, 14). To assess whether Cp can generate an isoDGR motif able to recognize integrins in physiological conformations, we analyzed the effect of Cp aged under oxidative conditions on the adhesion of human epithelial (HaCaT), endothelial (E.A.hy926), and glioblastoma (T98G) cell lines. Preliminary flow cytometry analysis showed αvβ6 expression by HaCaT, αvβ3 and αvβ5 expression by EA.hy926, and αvβ8 expression by T98G (data not shown).

Plates coated with ox/Ambic-Cp increased in a dose-dependent manner the adhesion and spreading of HaCaT cells, whereas untreated Cp induced little cell adhesion (Fig. 5A and B). In HaCat cells adhering to ox/ambic-Cp coated plates, cell spreading showed a significant (p = 0.0087) area increase (~20%) as compared with cells adhering to Cp coated plates (Fig. 5C). Similar pro-adhesive effects were observed with glioblastoma (Fig. 5D and E) and endothelial cell lines (data not shown).

To assess whether cell adhesion was mediated by isoDGR, we incubated ox/Ambic-treated Cp with PIMT, an enzyme that converts L-isoAsp residues to L-Asp (10, 30). PIMT treatment almost completely inhibited the pro-adhesive activity of ox/Ambic-treated Cp and of isoDGR peptide, the latter serving as a control (Student’s t test, p < 0.0001) (Fig. 5D and E). These results suggest that isoAsp formation, presumably at NGR site(s) of Cp, is associated with a “gain of function” in cell adhesion assays.

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9, and 12 days at 37 °C. Cp was added in large amounts because of the presence of other integrin-binding molecules in CSF (e.g., fibronectin, tenascin) that can compete the binding of Cp. Aging in a pathological milieu was able to induce a time-dependent Cp significant binding to \( /H9251 /H9252 /H11021 /H0.0001 \) (Fig. 8 \( A \), CSF/AD), whereas binding was absent or very weak when aging occurred in normal CSF (Fig. 8 \( A \), CSF). Thus, Cp deamidation showed a faster kinetic in the CSF from AD patients than from healthy subjects. By comparison with the binding ability of the same amount of \( \text{in vitro} \) chemically aged Cp (Cp-ox/AmBic), we ascertained that \( \sim 40\% \) of the added Cp was converted to pro-adhesive Cp after 12 days of aging in AD patients CSF (Fig. 8\( B \)). Also in the case of purified Cp aging in the CSF, the gain in integrin binding properties was mediated by Cp deamidated NGR motifs, as demonstrated by competition with an isoDGR peptide, which abolished binding to \( /\alpha v /\beta 6 \) \((p < 0.0001)\) (Fig. 8\( C \), gray bars). To investigate whether Cp aged in pathological conditions was also able to recognize integrins in their physiological conformation, we performed cell adhesion experiments. HaCat cells were seeded on plates coated with Cp immunoprecipitated from either AD CSF or control CSF at time 0 or after 9 days of aging. The Cp aged in pathological CSF was able to promote the HaCat cell adhesion (Fig. 8D, black bars), whereas little or no adhesion was observed on Cp aged in control CSF (Fig. 8D, white bars). In addition to the acquisition of integrin binding function, Cp aged in AD CSF showed a complete loss of ferroxidase activity \((p < 0.0001)\), whereas the Cp aged in healthy CSF showed a reduction of \( \sim 40\% \) (Fig. 8E, 9 days).

DISCUSSION

Asparagine deamidation during protein aging depends on neighboring residues as well as on secondary and tertiary structural elements. The Asn deamidation rate is also affected by environmental factors such as pH, temperature, and ionic strength \((1, 2)\). The combination of these features controls the kinetic of deamidation reactions that can range from hours to years \((1, 2)\). Thus, for each protein it is important to assess whether specific Asn residues can undergo or not this post-translational modification. For example, we previously reported that the deamidation of an NGR site of fibronectin does not occur in the intact protein, whereas it can rapidly occur in protein fragments \((10)\). The results reported here show that deamidation of
the $^{568}$NGR site, which is exposed on the protein surface, does not require structural changes. In contrast, deamidation of the $^{962}$NGR site, which is buried inside two $\beta$-strands of the protein structure and is blocked in a stable conformation by polar interactions with neighboring amino acids, can occur only after structural changes induced by an oxidative microenvironment.

Protein oxidation and conformational changes similar to those reported here have been shown to cause copper ions to release from Cp (17, 26, 28). Noticeably, copper ions are necessary for the Cp ferroxidase activity (17, 26, 28). Because protein aging under oxidative stress is a condition that may occur in some neurodegenerative diseases (19, 31–33), we hypothesize that Cp aging under pathological conditions promotes modifications that favor structural changes that in turn foster both copper ion release and NGR deamidation. Copper ions released from Cp might contribute to the reported increase of the pool of free copper and to the decrease of active Cp found in the CSF of AD patients (34–36). It is conceivable that this increase, affecting the redox environment (37), might contribute to the observed Cp modifications by a feedback mechanism. The observed loss of ferroxidase activity of Cp aged in AD CSF suggests that changes in the Cp structure can be fostered also ex vivo. Given the role of Cp, ferroxidase activity reduction may affect the iron homeostasis (38), causing an increase in iron retention as seen in (17, 26, 28).

A relationship between oxidation and protein deamidation in neurodegenerative disease is also suggested by the observation that several proteins found to be deamidated in the brain of PIMT knock-out mouse were oxidized in the brain of AD patients (30). We previously reported that in the CSF from Par-

FIGURE 7. Schematic representation of the signaling transduction throughout integrin engagement by Cp aged under oxidative conditions. Changes in molecule phosphorylation after 30 and 120 min of treatment with deamidated Cp suggest that early signal is addressed to gene activation, cell cycle induction, and MAPK signaling pathway activation, whereas late signal sustains actin cytoskeleton rearrangement, and cell cycle and proliferation arrest. The schematic evaluation is based on the results obtained by activation signal analysis performed with reverse phase protein arrays (Fig. 6) and reports the rate of phosphorylation compared with control conditions consisting in cells treated with nonmodified Cp.
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**A**  
CSF spiked with Cp (20 μg/ml)  
CSF spiked with Cp (20 μg/ml)  
CSF/AD  
CSF/AD  

**B**  
CSF spiked with Cp (20 μg/ml)  
CSF spiked with Cp (20 μg/ml)  
CSF/AD  
CSF/AD  

**C**  
isoDGR peptide (μg/ml)  
isoDGR peptide (μg/ml)  
isoDGR peptide (μg/ml)  
isoDGR peptide (μg/ml)  

**D**  
HaCat cell adhesion to plates coated with immunoprecipitated purified Cp (5 μg/ml)  
HaCat cell adhesion to plates coated with immunoprecipitated purified Cp (5 μg/ml)  
HaCat cell adhesion to plates coated with immunoprecipitated purified Cp (5 μg/ml)  
HaCat cell adhesion to plates coated with immunoprecipitated purified Cp (5 μg/ml)  

**E**  
Ferric oxidase activity of Cp purified from CSF (% of control)  
Ferric oxidase activity of Cp purified from CSF (% of control)  
Ferric oxidase activity of Cp purified from CSF (% of control)  
Ferric oxidase activity of Cp purified from CSF (% of control)  

**FIGURE 8.** Cerebrospinal fluid from Alzheimer disease patients fosters NGR motif deamidation and binding to αvβ6 integrin of spiked purified Cp. A, binding to αvβ6 of purified Cp added to CSF from healthy subjects (CSF, white squares) or AD patients (CSF/AD, black squares) aged for 0, 3, 6, 9, and 12 days at 37 °C. B, quantitation of the Cp bound to αvβ6 integrin after ex vivo aging in CSF evaluated by the comparison with binding of chemically in vitro aged Cp (Cp-ox/Ambic, 100%). Aging from 0 to 9 days at 37 °C of purified Cp in the CSF from AD patients (CSF/AD) showed ~40% of the protein binding ability to αvβ6, whereas the same aging conditions in the CSF from healthy subjects showed negligible binding ability (3–4%). C, competition with isoDGR peptide (gray bars) of the binding to αvβ6 of Cp aged in CSF from healthy subjects or AD patients (CSF and CSF/AD, respectively, white bars). D, HaCat cell adhesion to plates coated with immunoprecipitated purified Cp (5 μg/ml) after ex vivo aging (0 or 9 days) in CSF from healthy subjects (CSF, white bars) or AD patients (CSF/AD, black bars). Adhesion was evaluated as absorbance (570 nm) of the crystal violet-stained cells and was expressed as a percentage of control (cells in adhesion on BSA-coated wells). E, Cp ferric oxidase activity after incubation in CSF from healthy subjects (CSF, white bars) or AD patients (CSF/AD, black bars). Ferric oxidase activity of immunoprecipitated Cp (1.25 μg) was evaluated measuring the decrease in Bp-Fe(III) complex absorbance at 535 nm. The values are expressed as percentages of total ferric oxidase activity measured for 1.25 μg of untreated Cp. All the assays were performed in triplicate for two independent experiments, by using CSF from n = 8 different subjects each group (Table 1). Statistical p value was evaluated by Student’s t test. ***p < 0.0001; **p < 0.001; *p < 0.01.

Kinship and AD patients, Cp showed modifications derived from protein oxidation (17). Similar modifications have been reported for serum Cp during human aging (26). Thus, the Cp alterations observed in Parkinson disease and AD patients might reflect accelerated protein aging, as a consequence of changes in the environmental redox status of pathological CSF (18, 19). Therefore, it is plausible that the reported Cp oxidation-induced changes (17) also include Asn deamidation of NGR motifs.

Another major finding of this work is that the NGR to isoDGR conversion in Cp induces a gain of function in terms of binding to several αv integrins and, above all, the fact that the interaction of deamidated Cp with integrins can trigger an intracellular signaling cascade. We previously demonstrated that the NGR to isoDGR transition in fibronectin can work as a molecular switch for integrin-ligand recognition; the same mechanism seems to explain the gain of integrin binding function in Cp aged under oxidative conditions (10–12, 14). It is of relevance that, in contrast to fibronectin, the binding to integrin occurs for the intact full-length deamidated Cp. Consistently with binding data, our docking studies show that at least the 568iskoDGR motif of intact Cp can interact with the canonical RGD binding pocket of the αvβ6 having the stereochemical and electrostatic requirements for a correct recognition. Because it is not possible to predict a priori the putative structural rearrangement induced by the oxidation, we cannot describe in silico the possible interactions occurring between integrin and the 962iskoDGR sequence in the context of full-length Cp. The capability of the intact deamidated Cp to bind integrins might explain the ability to promote integrin activation.

These findings suggest that the Cp deamidation can produce a biological relevant gain of function. The functional importance of the NGR sites of Cp is also suggested by the high conservation of these tripeptide sequences across differing species from hamster to human. Remarkably, very few substitutions in various species were also shown in the 962NGR flanking sequences, which are likely crucial for regulating the deamidation rate.

Notably, Cp NGR deamidation might occur in a relevant manner in certain pathological conditions, as suggested by the observation that Cp ex vivo aged in the CSF from AD patients acquires integrin binding and cell adhesion properties. The faster kinetic of Cp deamidation observed in CSF from AD patients compared with the CSF from healthy subjects supports the hypothesis of a generally accelerated protein aging in neurodegenerative diseases, as a consequence of pathological environment (18, 19). Based on the observation that after 12 days of aging in the pathological CSF ~40% of the Cp were able to bind integrin and given that the Cp concentration in the CSF is ~2 μg/ml and that its physiological half-life is ~5.5 days in vivo (15), it is conceivable that NGR deamidation might occur in patients to an extent that yields functionally relevant concentration of deamidated Cp (~200 ng/ml). The observed induction of NGR modification in Cp incubated in the CSF from AD patients implies the presence of factors that foster/accelerate deamidation, either directly or indirectly as a consequence of the induction of structural changes. However, further studies are necessary to define the differential composition of healthy and AD CSFs as to identify factors underlying the increased deamidation rate observed in pathological CSF.
It is known that integrin binding may activate intracellular signaling cascades, which in turn can affect cell differentiation, survival, growth, and division (39). Thereby, the hypothesis that in neurodegenerative diseases Cp might deamidate and transduce unusual intracellular signals opens new investigation perspectives in the AD field. It has been reported that an aberrant signaling mediated via integrin engagement (e.g., αββ1, αββ3, and αββ5) by fibrillar amyloid-β in the CNS can result in unconventional FAK activation and downstream signaling (i.e., MAPK, GSKβ3, etc.) (40–42). A similar signaling mechanism is triggered by deamidated Cp in epithelial cells: we observed inhibition of FAK and MAPK signaling, which could lead to cell cycle arrest, proliferation inhibition, and cytoskeleton reorganization. Various cell types in the CNS might be activated via integrins, including microglial cells, which contribute to the inflammation mechanisms in neurodegenerative diseases (43). Interestingly, it has been reported that Cp can activate microglial cells (44), although the receptor underlying this mechanism has not yet been identified: one possibility is that deamidated Cp interacts with integrins. Other potential cell targets of deamidated Cp include the specialized epithelial cells of the ependymal layer and choroid plexus, which are directly in contact with the CSF and have been reported to be altered in AD (45–47).

In conclusion, the results show that Asn deamidation reactions, which may occur in Cp upon aging, can convert this protein into an isoDGR-containing ligand of αv integrins. Furthermore, integrin-mediated intracellular signaling can be transduced by the modified protein. Oxidation-induced structural changes foster the NGR to isoDGR transition at the NGR site buried in the protein structure. Even though the mechanism we described results mainly from in vitro observations, the ex vivo evidence that the CSF from AD patients promotes Cp deamidation suggests that Cp modifications might also occur in patients. Further investigations are needed to assess the biological role of in vivo Cp deamidation and the consequences for neurodegenerative diseases.

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