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A Residue-specific Shift in Stability and Amyloidogenicity of Antibody Variable Domains*

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Variable (V) domains of antibodies are essential for antigen recognition by our adaptive immune system. However, some variants of the light chain V domains (VL) form pathogenic amyloid fibrils in patients. It is far unclear which residues play a key role in governing these processes. Here, we show that the conserved residue 2 of VL domains is crucial for controlling its thermodynamic stability and fibril formation. Hydrophobic side chains at position 2 stabilize the domain, whereas charged residues destabilize and lead to amyloid fibril formation. NMR experiments identified several segments within the core of the VL domain to be affected by changes in residue 2. Furthermore, molecular dynamic simulations showed that hydrophobic side chains at position 2 remain buried in a hydrophobic pocket, and charged side chains show a high flexibility. This results in a predicted difference in the dissociation free energy of ~10 kJ mol⁻¹, which is in excellent agreement with our experimental values. Interestingly, this switch point is found only in VL domains of the κ family and not in VL domains, despite a highly similar domain architecture. Our results reveal novel insight into the architecture of variable domains and the prerequisites for formation of amyloid fibrils. This might also contribute to the rational design of stable variable antibody domains.

Antibodies consist of structurally highly homologous domains all sharing the immunoglobulin fold. This tertiary structure is characterized by two β-sheets that are connected by a conserved disulfide bridge and form a β-barrel structure. In the variable domains, the anti-parallel β-pleated sheets include nine β-strands in a 4 + 5 orientation (ABE(D/C)C′C′′FG) (1), whereas those of the constant domains are made up of seven β-strands in a 4 + 3 orientation (ABE(D/C)FG) (2). In the variable domains, this framework structure is expanded by three variable loops (CDR³ 1–3) that mediate antigen recognition. The principles that govern the stability of the variable domains are of special importance, as loops of different lengths and composition have to be tolerated. To achieve this, the amino acid composition of the framework is highly conserved at most positions. Single substitutions in variable domains can have deleterious effects, seen in some patients with monoclonal gammopathy or clonal B cell dyscrasias. In these patients, antibody variants are pathologically deposited in a fatal illness called light chain deposition disease (3, 4) and systemic amyloid light chain (AL) amyloidosis (5, 6). Analysis of proteins isolated from AL patient biopsies showed that the main constituent of amyloid deposits is the variable domain of the light chain (VL) (7, 8). The basis of the disease is the overproduction of unpaired antibody light chains (LCs), which are secreted into the bloodstream and taken up by multiple organs and tissues where they aggregate, severely compromising the functions of the affected organs and ultimately leading to death (9, 10). AL amyloidosis remains the most common and most fatal systemic amyloidosis in western countries (7, 11).

Despite the high similarity between the VL and VH domains, VH is less susceptible to amyloid fibril formation (12). Furthermore, why some VLs form amyloids and others do not still remains largely enigmatic. In general, it is therefore not clear which residues are of special importance for the structural integrity of the VL domain. As the C region domain is not involved in the deposits, it seems that structural elements exist that protect it from misfolding (13), whereas some factors within the VL domain contribute to the rational design of stable variable antibody domains.

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† To whom correspondence should be addressed. Tel.: 49-89-289-13340; Fax: 49-89-289-13345; E-mail: johannes.buchner@tum.de.
‡ The abbreviations used are: CDR, complementarity-determining region; AL, antibody light chain; TEM, transmission electron microscopy; LC, light chain; MD, molecular dynamics; r.m.s.d., root mean square deviation; PDB, Protein Data Bank; GdmCl, guanidinium chloride; US, umbrella sampling; ANS, 1-anilino-8-naphthalene sulfonate; V, variable; ThT, thioflavin T.
domain might predispose it to the disease state. Highly conserved consensus sequences within the framework region are potentially important elements in this respect. A striking feature of variable domains is the conserved N terminus of \( \beta \)-strand A (FWR1). However, the role of the N-terminal residues of the LC on domain structure, stability, and amyloidogenic potential is still unresolved.

In this study, we performed a detailed analysis of the importance of the conserved N-terminal residues for the structural integrity, the conformational stability, and amyloidogenicity of variable domains. With biophysical approaches including CD, fluorescence, NMR, and molecular dynamics (MD) simulations, we demonstrate that specifically residue 2 affects the conformational integrity and is a key element controlling its amyloidogenic properties. This reveals novel insights into the architecture of antibody variable domains.

**EXPERIMENTAL PROCEDURES**

Oligonucleotides (primers) were purchased from MWG Biotech. Utp1apure GdmCl was obtained from Sigma. All other chemicals were purchased from Merck. The concentrations of GdmCl solutions were determined from refractive indices (14). Unless stated otherwise, all measurements were carried out at 25 °C (pH 7.4) in PBS buffer.

**Cloning, Mutagenesis, Expression, and Purification of the Variable Domain Variants**—The MAK33 V\(_{1}\)\(_{\kappa}\) wild type encoding plasmid was obtained as published previously (15). 1OPG V\(_{1}\)\(_{\kappa}\), 1AQK V\(_{1}\)\(_{\lambda}\), and 1VGE V\(_{1}\)\(_{\lambda}\) wild type sequences cloned into the pET28b vector between the Ncol and HindIII restriction sites were purchased from Geneart AG (Regensburg, Germany). Single, double, and triple point mutants were generated with QuikChange mutagenesis PCR with forward and reverse primers carrying the target mutation. Inserts of the wild type, single point, and double point mutants in the pET28b vector were used as PCR templates for the generation of single, double, and triple point mutants, respectively. The PCR was followed by digestion of template DNA with DpnI enzyme (New England Biolabs) for 1 h at 37 °C. The PCR product was then transformed into E. coli Mach1 cells. Positive clones were verified by PCR and sequencing primer by Eurofins MWG Operon to verify DNA from a single colony was isolated with the Wizard Plus SV Miniprep kit (Promega) and sequenced using the T7 forward or reverse strand primer by Eurofins MWG Operon to verify.

**Growth, and inclusion bodies were prepared as described previously** (15, 16). In brief, the plasmid was transformed into E. coli BL21(DE3)-star cells (for V\(_{1}\)\(_{\kappa}\) and \( \lambda \) variants) or in E. coli JM109 cells (for V\(_{1}\)\(_{\kappa}\) variants) for expression at 37 °C. At an \( A_{\text{600}} \) of 0.6–0.8, expression was induced using 1 mM isopropyl \( \beta \)-D-thiogalactopyranoside. Cells were harvested after overnight growth, and inclusion bodies were prepared as described previously (17, 18). The pellet was solubilized and unfolded in 25 mM Tris-HCl (pH 8), 5 mM EDTA, 8 M urea, and 2 mM \( \beta \)-mercaptoethanol at room temperature for at least 2 h. The soluble fraction was then injected onto a Q-Sepharose column equilibrated in 25 mM Tris-HCl (pH 8), 5 mM EDTA, and 5 M urea. The proteins were eluted in the flow-through and diluted two times before being refolded by dialysis into 250 mM Tris-HCl (pH 8.0), 100 mM L-Arg (for V\(_{1}\)\(_{\kappa}\) and \( \lambda \) variants) or 400 mM L-Arg (for V\(_{1}\)\(_{\lambda}\) variants), 5 mM EDTA, 1 mM oxidized glutathione, and 0.5 mM reduced glutathione at 4 °C overnight. To remove misfolded aggregates and remaining impurities, the protein was cleaned using a Superdex 75 gel filtration column (GE Healthcare) equilibrated in PBS buffer. The recovery and purity of intact protein were verified by SDS-PAGE and matrix-assisted laser desorption/ionization time-of-flight mass spectrometry.

**CD Measurements**—CD measurements were performed using a Jasco J-720 spectropolarimeter (Jasco, Grossumstadt, Germany) equipped with a Peltier element. Far-UV CD spectra were measured using 10 \( \mu \)M protein in a 1-mm path length cuvette between 260 and 200 nm. Near-UV CD was measured between 320 and 250 nm using 50 \( \mu \)M protein in a 1-mm cuvette. All spectra were accumulated 16 times, and buffer was corrected. Thermal transitions were recorded using 10 \( \mu \)M protein at 212 nm for V\(_{1}\)\(_{\kappa}\) and 205 nm for V\(_{1}\)\(_{\lambda}\) and V\(_{1}\)\(_{\lambda}\) variants with a heating and cooling rate of 20 °C/h.

**Tryptophan Fluorescence Measurements**—Unless otherwise stated, all fluorescence measurements were carried out using a FluoroMax-4 spectrophuorimeter (Horioba Jobin Yvon, Bensheim, Germany). Tryptophan fluorescence measurements were performed with excitation and emission slit widths of 2 and 3 nm, respectively. The protein concentration in a 1-cm quartz cuvette was 1 \( \mu \)M. The sample was excited at 280 nm, and spectra were recorded between 300 and 450 nm. Equilibrium unfolding and refolding transitions were carried out by denaturing 1 \( \mu \)M protein overnight at different concentrations of GdmCl (between 0 and 3 M GdmCl). Fluorescence intensity at 358 nm was measured for 50 s, and the average was taken. Analysis of data was carried out using the linear extrapolation method described previously (14, 19).

**1-Arginine-8-napthalene Sulfonate (ANS) Binding Assay**—ANS binding was measured by fluorescence emission recorded from 400 to 650 nm, with excitation at 380 nm with excitation and emission slit widths of 2 and 3 nm, respectively. Experiments were performed at a protein concentration of 10 \( \mu \)M and an ANS concentration of 100 \( \mu \)M in 1-cm quartz cuvette. All spectra were accumulated three times and averaged.

**Fibril Formation with Ultrasoundation**—For the standard reaction, the V\(_{1}\)\(_{\kappa}\) variants were diluted at a concentration of 30 \( \mu \)M in PBS solution and 5 \( \mu \)M thioflavin T (ThT). Then 0.2 ml of the V\(_{1}\)\(_{\kappa}\) variant solution was applied to each of the 96 wells of a microplate (Greiner Bio-One on a 96-well microplate 675-074 made of polystyrene with a size of 128 × 85 mm). The microplate was set on the HANABI (HANdai Amyloid Burst Inducer), which combines the use of a water bath-type ultrasonicator (ELESTEIN, Elekon, Chiba, Japan) and a microplate reader (SH-9000, Corona Electric Co., Ibaraki, Japan). The water bath with 400 × 250 mm × 230 mm (height) contains about 12 liters of water, and three ultrasonic transducers are set, two on the sides and one at the bottom. Pulses were applied to the microplate from three directions, focusing on the center position, for cycles of 1 min followed by a quiescent period of 9 min. The measurement temperature was maintained at 37 °C. The formation of fibrils was monitored by ThT fluorescence.
with excitation and emission wavelengths of 450 and 490 nm, respectively. Microplates were incubated at 37 °C and shaken before the measurements. Transmission electron microscopy (TEM) images were taken using a HITACHI H-7650 transmission microscope, (Hitachi, Tokyo, Japan), with an acceleration voltage of 80 kV. Aliquots (5 μl) of the samples were placed on a 400-mesh copper grid covered by a carbon-coated colloidal film for 60 s. Two percent (w/v) uranyl acetate solution (5 μl) was then placed on the grid to stain the samples for 60 s. Excess sample solutions on the grid were removed with a filter paper. The magnification on the representative pictures was 30,000-fold.

Fibril Formation without Ultrasonication—Samples of VΔ at a concentration of 30 μM in PBS buffer (pH 7.4), containing 0.05% sodium azide (or acetate buffer (pH 2)) were incubated with shaking (gentle agitation) in a roller shaker at 37 °C for at least 1 week. At various intervals, a 20-μl aliquot was withdrawn for ThT binding and TEM. For the ThT binding, 10 μl of protein aliquot was added to 480 μl of a PBS buffer (pH 7.4) followed by the addition of 10 μl of 500 μM ThT prepared in PBS buffer (pH 7.4) (ThT final concentration 10 μM). This was left to incubate for 90 s before acquiring the emitted signal in a 1-cm quartz cuvette from 430 to 570 nm with excitation at 440 nm. The spectra were recorded using an FP-6500 spectrofluorimeter (Jasco) equipped with a Jasco ADP-303T Peltier as temperature controller. The excitation and emission slit widths were both 3 nm. A scan speed of 100 nm/min, an integration time of 1 s, and a 0.5-nm recording interval were used. Three scans were averaged for each sample. Rayleigh (elastic) light scattering at an angle of 90° was also measured at the excitation wavelength peak of 440 nm (20, 21). Therefore, for the same sample, information on its light scattering ability (presence of macromolecular assemblies or aggregation) and ThT binding (fibril formation) was assessed. For comparison, the Rayleigh scattering intensity at 440 nm and the ThT fluorescence intensity at 480 nm were used.

For TEM measurements, a 10-μl protein aliquot was fixed on a 200-mesh activated copper grid and incubated for 60 s. After a washing step with 10 μl of H2O, the samples were stained with 10 μl of a 1.5% uranyl acetate solution. Samples were recorded with an acceleration voltage of 100 kV at a magnification of ×33,000 on a transmission electron microscope JEM100CX (jeol).

Thermally Induced Aggregation—Native MAK33 or 10PG VΔ at a concentration of 10 μM in PBS buffer (pH 7.4) with 10 μM ThT was subjected to increasing temperatures from 20 to 80 °C in steps of 5 °C in a 1-cm path cuvette. After thermal equilibration for 3 min, emission spectra were recorded for Rayleigh scattering and ThT fluorescence (21) as stated under “Fibril Formation without Ultrasonication.”

NMR Spectroscopy—All NMR spectra were recorded using uniformly 15N,13C-labeled proteins in 20 mM phosphate, 50 mM NaCl (pH 6.5) at 298 K. Backbone resonance assignments of MAK33 VΔ WT and MAK33 VΔ were transferred from previous assignments of MAK33 VΔ (data not shown). The assignment was accomplished based on three-dimensional HNCA, HCACAB, CBCA(CONH), HNCO, and HNCACO experiments (22). Experiments were carried out at 1 mM protein concentration. Those residues, for which the assignment could not be transferred from MAK33 VΔ-S20N, were assigned based on an HNCA experiment using a 100 μM MAK33 VΔ-12E sample. All spectra were recorded on a Bruker AVANCE 900MHz spectrometer (Bruker Biospin) equipped with a triple-resonance cryoprobe. Spectra were processed using TOPSPIN 3.2 (Bruker BioSpin) and analyzed with CcpNmr 2.2.2 (23). For determination of residues affected by the 12E substitution, 15N HSQC spectra of MAK33 VΔ WT and MAK33 VΔ-12E, both at 50 μM, were acquired on a Bruker Avance 600MHz spectrometer equipped with a cryoprobe. For all residues, chemical shift perturbations were determined using Equation 1,

\[ \Delta \delta^{res} = \sqrt{\left(\Delta \delta^{1H}\right)^2 + \frac{1}{25} \left(\Delta \delta^{15N}\right)^2} \]

with \( \Delta \delta^{res} \) being the weighted chemical shift difference, \( \Delta \delta^{1H} \) as the chemical shift of the amide proton in ppm, and \( \Delta \delta^{15N} \) as the chemical shift of the amide nitrogen in ppm.

For comparison of signal intensities, all signals, which displayed significant overlap in one of the proteins, were ignored. Intensities were normalized for both proteins to the same average intensity.

MD and Umbrella Sampling (US) Simulations—Start structures of MAK33 VΔ, κ domains and VΔ, α and VΔ, H domains were obtained by extracting the corresponding coordinates from the crystal structures 1FH5 (MAK33 VΔ, κ), 10PG, 1AQK, and 1VGE. Residue 2 of MAK33 VΔ, κ was substituted with Glu although residue 2 of 10PG, and VΔ, κ was replaced with Ile. In silico substitutions were performed using the SPDBV package (24), while selecting the best fitting side chain rotamer. All MD simulations and the analysis of root mean square deviations (r.m.s.d.) and fluctuations were performed using the Amber12 package (25). Proteins were solvated in octahedral boxes, including explicit ions and explicit (TIP3P) water molecules (26). The simulation systems were first energy-minimized (5000 steps) followed by heating up to 300 K in steps of 100 K with position restraints on all heavy atoms of the proteins. Subsequently, positional restraints were gradually removed from an initial 25 kcal mol\(^{-1}\) Å\(^{-2}\) to 0.5 kcal mol\(^{-1}\) Å\(^{-2}\) within 0.5 ns followed by a 1-ns unrestrained equilibration at 300 K. All production simulations were performed at a temperature of 300 K and a pressure of 1 bar. US simulations were performed using the distance between the Cα atom of residue 2 and the Cα of residue 32 at the floor of the binding region for residue 2 in the VΔ, κ domains as a reaction coordinate. A quadratic penalty potential \( k (d_{Cα-Cα} - d_{ref})^2 \), force constant \( k = 2.0 \) kcal mol\(^{-1}\) Å\(^{-2}\) for the Cα-Cα distance was used with reference distances varying from 11.5 to 16 Å in 0.5-Å steps and from 16 to 20 Å in 1-Å steps. At ~12–13 Å, residue 2 stays bound to the protein in the cavity as observed in the experimental x-ray structure, whereas it adopts a fully exposed state at distances of >16 Å. The associated potential of mean force was calculated using the weighted histogram analysis method (27).

Role of Residue 2 for Integrity of Antibody Variable Domains
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RESULTS

Biophysical Characteristics, Stability, and Amyloidogenic Propensity of Two Highly Homologous $\gamma$/k Domains—MAK33 is a well studied IgG antibody with respect to folding and association. In this context, the folding pathway of the $\gamma$/k domain has been analyzed in detail (15); its amino acid composition is typical for a murine $\gamma$/k domain and characterized. The far-UV circular dichroism (CD) spectra of MAK33 $\gamma$/k and 1OPG $\gamma$/k, already a partial exposure of tryptophan in the native 1OPG domain, were monitored by Rayleigh (elastic) light scattering at 440 nm. Their near-UV CD spectra (Fig. 1a) with a minimum at 275 nm confirmed that the tertiary structure of both proteins is correctly formed (30, 31). The presence of a buried tryptophan in the disulfide in antibody domains makes it a sensitive probe for studying conformational changes by fluorescence. Intrinsic tryptophan fluorescence spectra of native MAK33 $\gamma$/k and 1OPG $\gamma$/k domains monitored by Rayleigh (elastic) light scattering at 440 nm.

FIGURE 1. Sequence analysis of $\gamma$/k domains, spectroscopic characterization of MAK33 $\gamma$/k and 1OPG $\gamma$/k. a, multiple sequence alignment of $\gamma$/k sequences. Five representative sequences are shown. MAK33 $\gamma$/k differs from 1VGE $\gamma$/k at 40 positions and from 1OPG $\gamma$/k at five positions as marked by red rectangles. Identical CDRs between MAK33 $\gamma$/k and 1OPG $\gamma$/k domains are highlighted in different colors. b, far-UV CD spectra of native (continuous line) and temperature-denatured (dotted line; 60 °C) MAK33 $\gamma$/k and 1OPG $\gamma$/k in PBS buffer. c, near-UV CD spectra of native MAK33 $\gamma$/k (green line) and 1OPG $\gamma$/k (blue line). Intrinsic tryptophan fluorescence spectra of native (continuous line) and 3 M GdmCl-denatured (dotted lines) MAK3 3 $\gamma$/k (d) and 1OPG $\gamma$/k (e) in PBS buffer. f, ANS fluorescence spectra of native MAK33 $\gamma$/k (green line) and 1OPG $\gamma$/k (blue line). The gray line is PBS/ANS without protein. g, thermally induced aggregation of MAK33 $\gamma$/k and 1OPG $\gamma$/k domains monitored by Rayleigh (elastic) light scattering at 440 nm.

random coils, which might be due to long flexible CDR loops. Their near-UV CD spectra (Fig. 1c) with a minimum at 275 nm confirmed that the tertiary structure of both proteins is correctly formed (30, 31). The presence of a buried tryptophan in close proximity to the disulfide in antibody domains makes it a sensitive probe for studying conformational changes by fluorescence. Intrinsic tryptophan fluorescence spectra of both $\gamma$/k domains showed MAK33 $\gamma$/k to have a large (~2.5-fold) increase in fluorescence emission intensity (maximum at ~358 nm) in the presence of 3 M GdmCl (Fig. 1d), as opposed to a small increase observed in 1OPG $\gamma$/k (Fig. 1e). This suggests already a partial exposure of tryptophan in the native 1OPG $\gamma$/k. To verify whether the $\gamma$/k domains exposed hydrophobic patches on their surfaces, ANS binding was performed. As depicted in Fig. 1f, 1OPG $\gamma$/k binds ANS with an increase and a blue-shifted fluorescence emission maximum (shift from 525
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The propensity of both MAK33 and 1OPG V_{1,κ} domains to aggregate with increasing temperatures was monitored by recording Rayleigh (elastic) scattering of ThT fluorescence excitation (440 nm) light (Fig. 1f) (20, 21). Surprisingly, 1OPG V_{1,κ} already starts aggregating at ~35 °C, although MAK33 V_{1,κ} only starts transforming into aggregates at ~50 °C (a difference of ~15 °C). Maximum aggregation for both proteins occurred at ~65 °C. No ThT bonding was observed for any of the proteins at the experimental temperature range of 20–80 °C (data not shown), which implies that the aggregates formed by thermal induction did not contain amyloid fibrils. When we determined their stabilities against temperature monitored by far UV CD, we found, interestingly, that the 1OPG V_{1,κ} is 16 °C less stable than MAK33 V_{1,κ} as judged from their transition midpoints (Fig. 2b), which correlates very well with thermal-induced aggregation monitored by light scattering at 440 nm. Thermally induced unfolding transitions were not reversible for any of the proteins. For equilibrium unfolding and refolding in the presence of GdmCl monitored by Trp fluorescence, the MAK33 V_{1,κ} with a cooperative sigmoidal transition had identical midpoints in both the unfolding and refolding directions (Fig. 2c).

![FIGURE 2. Stability and amyloidogenic propensity of MAK33 V_{1,κ} and 1OPG V_{1,κ} domains.](http://www.jbc.org/

- **a**, thermal unfolding transitions of MAK33 V_{1,κ} (green symbols) and 1OPG V_{1,κ} (blue symbols). The solid lines indicate the theoretical curves derived by fitting the data to a Boltzmann function for MAK33 V_{1,κ} (green) and 1OPG V_{1,κ} (blue) to obtain transition midpoints (T_m). **b**, GdmCl-induced unfolding transitions of MAK33 V_{1,κ} (green) and 1OPG V_{1,κ} (blue). The reversibility of the GdmCl-induced unfolding process is shown by the overlay of unfolding symbols (open circles) and refolding (closed circles) experiments. The solid lines show the fit to a two-state mechanism for both MAK33 V_{1,κ} (green line) and 1OPG V_{1,κ} (blue line) to obtain thermodynamic stability values (ΔG_u) and the cooperativity parameters (m values). **c**, transmission electron microscopy micrographs of MAK33 V_{1,κ} and 1OPG V_{1,κ} from amyloid induction experiments at neutral pH, 37 °C, and 1 week incubation with gentle agitation. Scale bars, 200 nm.

![1OPG VLk-wt](http://www.jbc.org/)

![MAK33 VLk-wt](http://www.jbc.org/)

pH 7.4

to 515 nm), suggesting the presence of exposed hydrophobic patches. MAK33 V_{1,κ} shows no ANS binding (Fig. 1f).
Role of Residue 2 for Integrity of Antibody Variable Domains

![Residue frequency distribution analysis at position 2 of variable domains and of the different positions between MAK33 $\kappa$ and 1OPG $\kappa$ domains. Relative distribution of amino acid at various positions of light (a–e) and heavy chains (f). Green arrowheads indicate residue in MAK33 $\kappa$, and the corresponding positions in 1OPG $\kappa$ are shown by blue arrowheads. With the exception of positions 100 and 106, the residues found in MAK33 $\kappa$ at positions 2, 3, and 102 are more frequent than those at corresponding positions in 1OPG $\kappa$. The database could not separate $\lambda$ LCs from the $\kappa$ LCs during the distribution analysis. But an observation of most of the sequences show that Ser-2 is almost entirely contributed by the $\lambda$ LCs (orange arrowhead in a). f, red arrowheads indicate the most frequent residue at position 2 of heavy chains. Sequences from all organisms were considered for the analysis. Analysis was based on the Kabat numbering scheme accessed through the antibody (abYsis database) (Dr. Andrew C. R. Martin’s Group, University College London).](http://www.jbc.org/)

FIGURE 3. Residue frequency distribution analysis at position 2 of variable domains and of the different positions between MAK33 $\kappa$ and 1OPG $\kappa$ domains. Relative distribution of amino acid at various positions of light (a–e) and heavy chains (f). Green arrowheads indicate residue in MAK33 $\kappa$, and the corresponding positions in 1OPG $\kappa$ are shown by blue arrowheads. With the exception of positions 100 and 106, the residues found in MAK33 $\kappa$ at positions 2, 3, and 102 are more frequent than those at corresponding positions in 1OPG $\kappa$. The database could not separate $\lambda$ LCs from the $\kappa$ LCs during the distribution analysis. But an observation of most of the sequences show that Ser-2 is almost entirely contributed by the $\lambda$ LCs (orange arrowhead in a). f, red arrowheads indicate the most frequent residue at position 2 of heavy chains. Sequences from all organisms were considered for the analysis. Analysis was based on the Kabat numbering scheme accessed through the antibody (abYsis database) (Dr. Andrew C. R. Martin’s Group, University College London).

tively, which reflects the different cooperativities observed. Although the stability of MAK33 $\kappa$ falls within the range of those reported for amyloidogenic $\lambda$ domains (15–20 kJ mol$^{-1}$) (32, 33), it is known to be nonamyloidogenic in vitro at physiological pH (15). Accordingly, after 1 week of incubation in PBS buffer at pH 7.4 and 37 °C with gentle agitation, MAK33 $\kappa$ did not form any amyloid fibrils (Fig. 2c). However, under these conditions, 1OPG $\kappa$ readily turned into well defined fibrillar structures (Fig. 2c). When the proteins were incubated in an acidic buffer (pH 2) under similar conditions, fibrils were detected for both MAK33 $\kappa$ and 1OPG $\kappa$ samples (data not shown), confirming our previous results on MAK33 $\kappa$ (15). This supports the notion that under appropriate destabilizing conditions, even proteins not known to form amyloid fibrils under normal physiological conditions can indeed fibrillize (34–36).

To assess how conserved the frameworks of both $\kappa$ domains are in comparison with other $\kappa$ sequences, a comprehensive sequence analysis was performed using the abYsis database, which integrates sequences from the Kabat (37) and IMGT (38) databases. Fig. 1a depicts five representative $\kappa$ sequences from this alignment. Considering amino acid side chain chemistry, all framework residues are very similar, except for residue 2. In 1OPG $\kappa$, this residue is a Glu, although in most other $\kappa$ frameworks an aliphatic residue is found at the same position (Fig. 1a). A residue frequency distribution analysis using the abYsis database with all available (~20,000 non-identical) LC sequences revealed that Ile occurs with the highest frequency (46%) at position 2 (Fig. 3). Other aliphatic residues like Val, Leu, and Ala represent 7–9% at this position in $\lambda$ domains and charged residues (Glu, Asp, Lys, and Arg) only <1% (Fig. 3a). In addition, looking at the residue frequency distribution at position 102 of these $\lambda$ domains, we observed a highly conserved Thr with a frequency of 98%, although all other residues had a <1% frequency of occurrence (Fig. 3d). In the more conformationally stable MAK33 $\kappa$, position 102 is occupied by Thr, whereas in the less stable 1OPG $\kappa$, the less frequent Ser is found. With positions 2 and 102 showing huge frequency distribution discrepancies between the two $\kappa$ domains, we wondered whether these positions were responsible for the different behaviors of the two proteins. In consequence, we exchanged these residues between the two $\kappa$ domains and monitored their effects on stability and amyloid fibril formation.

Residue 2 Is Crucial for the Stability of $\lambda$ Domains—To identify residue(s) that predispose 1OPG $\kappa$ for lower stability and fibril formation, single point variants were generated by replacing residues in 1OPG $\kappa$ with corresponding residues of MAK33 $\lambda$, at any of the five different positions (OP-E2I, OP-L3V, OP-G100A, OP-S102T, and OP-I106L). All five substitutions were correctly folded as revealed by UV CD spectra. The thermal unfolding of these variants showed a striking increase in thermal stability for OP-E2I with a $T_{\text{mol}}$ value of 43.1 °C, compared with 31.6 °C for 1OPG $\kappa$-WT. Also, OP-E2I was significantly more stable toward GdmCl than 1OPG $\kappa$-WT (Fig. 4, a and c; Table 1). OP-S102T was only slightly more stable, and the other single point substitution variants were all similar to the less stable 1OPG $\kappa$-WT (Table
FIGURE 4. Stability of the different \( V_{\kappa} \) variants. Thermal unfolding transitions of 1OPG \( V_{\kappa} \) variants (a), MAK33 \( V_{\kappa} \) variants (b), and 1VGE \( V_{\kappa} \) variants (d). The solid lines indicate the theoretical curves derived by fitting the data to a Boltzmann function to obtain transition midpoints (\( T_{\text{melt}} \)). c, increase or decrease in stability caused by the different substitutions in 1OPG \( V_{\kappa} \) and MAK33 \( V_{\kappa} \), obtained by subtracting the \( T_{\text{melt}} \) and \( \Delta G \) values of the wild type \( V_{\kappa} \) from those of their corresponding mutants. d, GdmCl-induced unfolding transitions of 1VGE \( V_{\kappa} \) variants. The reversibility of the unfolding process is shown by the overlay of unfolding symbols (open circles) and refolding (closed circles) experiments. The solid lines show the fit to a two-state mechanism for \( V_{\kappa} \) variants to obtain thermodynamic stability values (\( \Delta G_{\text{m}} \)) and the cooperativity parameters (\( m \) values).
TABLE 1
Thermal and chemical stabilities of 1OPG VL, κ and MAK33 VL, κ and 1VGE VL, κ variants

<table>
<thead>
<tr>
<th>Protein variant</th>
<th>$T_{melt}$</th>
<th>$\Delta G_{U}$</th>
<th>$M$ value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>$^\circ C$</td>
<td>$k$J mol$^{-1}$</td>
<td>$k$J mol$^{-1}$</td>
</tr>
<tr>
<td>1OPG VL, κ-WT</td>
<td>31.6 ± 0.2</td>
<td>−3.0 ± 2.6</td>
<td>12.9 ± 5.3</td>
</tr>
<tr>
<td>OP-E2I</td>
<td>43.1 ± 0.1</td>
<td>−5.1 ± 1.7</td>
<td>14.2 ± 1.7</td>
</tr>
<tr>
<td>OP-S102T</td>
<td>34.0 ± 0.2</td>
<td>−6.0 ± 1.6</td>
<td>16.6 ± 3.3</td>
</tr>
<tr>
<td>OP-E2I/S102T</td>
<td>45.8 ± 0.2</td>
<td>−13.8 ± 2.9</td>
<td>18.1 ± 4.4</td>
</tr>
<tr>
<td>OP-E2I/G100A/S102T</td>
<td>44.8 ± 0.4</td>
<td>−14.8 ± 4.0</td>
<td>16.4 ± 4.0</td>
</tr>
<tr>
<td>MAK33 VL, κ-WT</td>
<td>47.7 ± 0.2</td>
<td>−17.8 ± 2.6</td>
<td>18.0 ± 2.4</td>
</tr>
<tr>
<td>MK-12E</td>
<td>38.2 ± 0.1</td>
<td>−5.8 ± 1.6</td>
<td>15.0 ± 3.0</td>
</tr>
<tr>
<td>MK-T102S</td>
<td>46.0 ± 0.2</td>
<td>−13.8 ± 3.1</td>
<td>18.2 ± 3.2</td>
</tr>
<tr>
<td>MK-I2E/T102S</td>
<td>33.6 ± 0.2</td>
<td>−8.0 ± 3.0</td>
<td>19.8 ± 2.3</td>
</tr>
<tr>
<td>MK-ΔD1</td>
<td>48.2 ± 0.2</td>
<td>−14.2 ± 3.7</td>
<td>13.5 ± 3.3</td>
</tr>
<tr>
<td>MK-ΔD1/Δ2</td>
<td>41.7 ± 0.2</td>
<td>−9.7 ± 1.7</td>
<td>15.6 ± 2.5</td>
</tr>
<tr>
<td>1VGE VL, κ-WT</td>
<td>46.0 ± 0.1</td>
<td>−3.3 ± 0.4</td>
<td>10.7 ± 0.5</td>
</tr>
<tr>
<td>1VGE VL, κ-I2E</td>
<td>38.6 ± 0.2</td>
<td>−4.8 ± 0.6</td>
<td>10.7 ± 1.0</td>
</tr>
</tbody>
</table>

1), implying that the exchange of these residues is not important for the conformational stability of the domain. With the high increase in stability seen in OP-E2I and the small increase observed in OP-S102T, we next tested whether a combination of both substitutions (OP-E2I/S102T) and a triple substitution OP-E2I/G100A/S102T (OP-IAT) would further lead to increased stability. The double and triple substitution variants both unfolded with a $T_{melt}$ of about 45 °C quite similar to that obtained for OP-E2I (Table 1), confirming only a small influence of the G100A and S102T substitutions on the thermal stability of the protein. However, in terms of stability to GdmCl, both variants were more stable than the respective single substitution variants (Fig. 4c; Table 1). These stability values observed for the double and triple substitution variants of 1OPG VL, κ are similar to that of MAK33 VL, κ wild type (Table 1). This demonstrates an additive effect of exchanges at different positions.

Because Glu-2 and Ser-102 to a minor extent were determined to be the destabilizing residues in 1OPG VL, κ, we were also interested in how they affect MAK33 VL, κ. Interestingly, MK-12E was destabilized, whereas MK-T102S had stability values similar to that of MAK33 VL, κ-WT (Fig. 4, b and c; Table 1). Specifically, MK-12E was ∼10 °C less stable with a free energy of −12 kJ mol$^{-1}$ less than that of MAK33 VL, κ-WT. The combined substitution variant (MK-12E/T102S) was less stable than both single substitution variants with a $T_{melt}$ of 33.6 °C and a chemical stability of −8.0 kJ mol$^{-1}$ (Fig. 4c; Table 1). This again emphasizes the additive effect of both positions. To demonstrate that our findings on the pivotal role of residue 2 are general to the VL, κ family, we investigated a human κ/IgG1 antibody VL, κ domain (1VGE VL, κ) (39), which differs greatly in amino acid sequence from 1OPG and MAK33 VL, κ. The N-terminal motif is ELV as opposed to DIV in MAK33 and DEL in 1OPG VL, κ (Fig. 1a). To assess the role of residue 2 in this human VL, κ, Leu was replaced with Glu (1VGE VL, κ-L2E). Notably, analysis of their thermal and thermodynamic stabilities also revealed a decrease in thermal stability of ∼10 °C and a 2-fold decrease in thermodynamic stability for 1VGE VL, κ-L2E compared with the wild type (Fig. 4, d and e; Table 1). This correlates very well with results obtained for MAK33 and 1OPG VL, κ, implying that the effects mediated by residue 2 are conserved within the VL, κ family.

Influence of Sequence Variation on Amyloid Fibril Formation—To examine whether the increase in the stability of 1OPG VL, κ variants also correlates with the resistance to amyloid formation, we set up ultrafiltration amyloid induction assays at neutral pH at 37 °C (40–45). ThT fluorescence was used to monitor the fibrillation process (46, 47). All variants that had a thermodynamic stability of <10.0 kJ mol$^{-1}$ (Table 1) bound ThT as indicated by an increase in fluorescence after a lag time of several hours, whereas the more stable variants (OP-E2I/S102T, OP-IAT, MAK33 VL, κ-WT, and MK-T102S) showed no ThT fluorescence (Fig. 5, a and b). The large fluctuations in ThT fluorescence amplitude are likely due to differences in higher order structure of amyloid fibrils as ThT fluorescence can change depending on the morphology of amyloid fibrils (41–43, 48). These variations in ThT fluorescence amplitudes might also depend on the mutants, and it is worth noting that ThT fluorescence is not always proportional to the amount of amyloid fibrils (48).

To confirm the presence of fibrils, transmission electron microscopy was performed. As expected, amyloid fibrils were detected in all variants for which ThT fluorescence was observed (Fig. 5c). Similar results were obtained with gentle agitation without ultrafiltration after 4 or 10 days of incubation of the VL, κ variants (data not shown). However, although the stable variants were protected from amyloid fibril formation, they did transform into amorphous aggregates upon prolonged incubation at 37 °C, revealed by the increase in Rayleigh scattering intensity (data not shown) Thus, the changes in $T_{melt}$ domain stability induced by modulating residue 2 inversely correlated with its ability to form amyloid fibrils.

Residue 2 Is a Decisive Factor for VL, κ Domain Architecture—Alignment of VL, κ sequences revealed a highly conserved N terminus for this family, with residue 1 always being an acidic residue (Asp or Glu) and residue 2 mostly an aliphatic residue (Ile, Leu, or Val) (Fig. 1a). We wondered whether the conserved residue 1 also affects the stability of VL, κ. Thus, N-terminal truncation variants of MAK33 VL, κ were generated in which either the first residue (Asp-1) was deleted (MK-VL, κΔD1) or one in which both the first (Asp-1) and the second (Ile-2) residues were truncated (MK-VL, κΔD1/Δ2). Deletion of these N-terminal residues did not influence the structure of the domain as determined by CD spectroscopy. Surprisingly, the deletion of Asp-1 (MK-VL, κΔD1) did not lead to changes in the thermodynamic stability when compared with the wild type domain. In contrast, the deletion of both Asp-1 and Ile-2 (MK-VL, κΔD1/Δ2) resulted in a decrease in $T_{melt}$ and thermodynamic stability of ∼7 °C and ∼10 kJ mol$^{-1}$, respectively (Fig. 6, a and b; Table 1), which is similar to the stability obtained by substituting Ile-2 alone with Glu (MK-κ-I2E). These results show that residue 2 and not residue 1 is important for the stability of VL, κ domains.

With Glu at position 2 identified to be destabilizing and Ile or Leu at the same position are stabilizing, we were interested to determine how residues with other side chains might affect the stability of these VL, κ domains. Considering the residue frequency distribution at position 2 of VL, κ (Fig. 3a), residue 2 of
1OPG V\(_1\kappa\)-WT and MAK33 V\(_1\kappa\)-WT were replaced with Ala, Leu, Asp, Lys, or Gln. Interestingly, the thermal unfolding transitions of these variants revealed striking differences. All variants with an uncharged residue at position 2 had a high thermal stability (Table 2). Although the respective 1OPG V\(_1\kappa\) variants showed an increase in \(T_{melt}\) of 4–12 °C compared with their

FIGURE 5. Amyloidogenicity of V\(_1\kappa\) variants. 30 μM of each V\(_1\kappa\) variant in a PBS buffer at pH 7.4 in the presence of ThT was subjected to ultrasonic pulses at 37 °C (n = 3). ThT fluorescence was monitored over time for 1OPG V\(_1\kappa\) variants (a) and MAK33 V\(_1\kappa\) variants (b), and TEM micrographs (c) were acquired at the end of the assay to detect the presence of fibrils. Variants with higher thermodynamic stabilities (>10 kJ mol\(^{-1}\)), OP-E2I/S102T and OP-IAT in a and MK-VL\(_1\kappa\)-WT and MK-T102S in b, did not bind ThT, and as a result the curves of both proteins are superimposed.
wild type, MAK33 VL/H9260 variants with the same uncharged residues at position 2 had Tmelt values similar to the more stable MAK33 VL/H9260-WT (Fig. 6e; Table 2). However, all variants with a charged residue (whether positive or negative) at position 2 were strongly destabilized. OP-E2D and OP-E2K were as unstable as 1OPG VL/H9260-WT, whereas the same substitutions in MAK33 VL/H9260 resulted in an ~10°C decrease in Tmelt (Fig. 6e; Table 2).

Residue 2 Does Not Affect the Stability of VL/H9261 and VH Domains—Irrespective of the family, variable domains (VL/H9260, VL/H9261, and VH) share a very similar topology of variable CDRs and...
We were interested in determining whether residue 2 also plays a similar role in the VH domain. A multiple sequence alignment (Fig. 7, a and b) and a residue frequency distribution analysis at position 2 of HL and VH showed this position to be mostly occupied by Ser or Ala (Fig. 3a), although at the same position in VH domains, Val occurs with a frequency of 83% (Fig. 3f). To ascertain the effects of residue 2 in these variable domains, a human IgG1 VL (1AQK (49)) and a human IgG1 VH (1VGE (39)) were examined. In the 1AQK VL domain, residue 2 (Asn-2) was replaced with either a Glu (1AQK VL-N2E) or Ser (1AQK VL-N2S). Unexpectedly, stability studies did not reveal any differences between the wild type and the 1AQK VL-N2E mutant (Fig. 7, c and d; Table 3). This suggests that the stabiliz-
ing effects of residue 2 seen in the V_{\text{L,\kappa}} family cannot be ascribed to the V_{\text{L,\alpha}} family. Moreover, the mutant that contained Ser at position 2 (1AQK V_{\text{L,\alpha}}-N2S) was slightly more stable than the wild type and 1AQK V_{\text{L,\alpha}}-N2E mutant. When Val-2 of 1VGE V_{\text{H}} was substituted with a Glu (1VGE V_{\text{H}}-V2E) or Ala (1VGE V_{\text{H}}-V2A), again no pronounced thermal stability differences were observed (Fig. 7e; Table 3). Both 1VGE V_{\text{H}}-V2E and 1VGE V_{\text{H}}-V2A mutants were $\sim$3 °C less stable than the wild type. Unlike in the V_{\text{L,\kappa}} family, the effects of residue 2 seen in V_{\text{L,\alpha}} are minor and not specific for any amino acid side chain tested.

### Influence of Residue 2 on Domain Structure and Dynamics—

To obtain a structural understanding of the influence of the residue at position 2 of V_{\text{L,\alpha}}, both MAK33 V_{\text{L,\kappa}}-WT and MAK33 V_{\text{L,\kappa}}-I2E were analyzed by solution-state NMR spectroscopy. 71% of the non-proline residues could be assigned (Fig. 8a). Backbone assignment was hampered by chemical exchange dynamics. The unassigned residues mainly include the regions from the N terminus to Pro-8, from His-41 to Tyr-50, and from Cys-88 to Gly-101. Fig. 8b depicts chemical shift changes caused by the I2E substitution. We find particularly large chemical shift differences for residues Cys-23 to Ser-40.

### TABLE 3

<table>
<thead>
<tr>
<th>Variable domain variant</th>
<th>$T_{\text{melt}}$</th>
<th>$\Delta G$</th>
<th>$M$ value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>°C</td>
<td>kJ mol$^{-1}$</td>
<td>kJ mol$^{-1}$ si$^{-1}$</td>
</tr>
<tr>
<td>1AQK V_{\text{L,\alpha}}-WT</td>
<td>44.0 ± 0.2</td>
<td>-5.7 ± 1.3</td>
<td>13.1 ± 1.1</td>
</tr>
<tr>
<td>AQ-N2E</td>
<td>44.9 ± 0.5</td>
<td>-6.4 ± 1.7</td>
<td>15.1 ± 1.9</td>
</tr>
<tr>
<td>AQ-N2S</td>
<td>44.9 ± 0.4</td>
<td>-11.1 ± 3.0</td>
<td>17.8 ± 3.6</td>
</tr>
<tr>
<td>1VGE-V_{\text{H}}-WT</td>
<td>46.5 ± 0.6</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>VG-V2E</td>
<td>43.1 ± 0.6</td>
<td>NA</td>
<td>NA</td>
</tr>
<tr>
<td>VG-V2A</td>
<td>43.8 ± 0.5</td>
<td>NA</td>
<td>NA</td>
</tr>
</tbody>
</table>

FIGURE 8. Structural properties of V_{\text{L,\alpha}} variants monitored by NMR spectroscopy. a, $^1$H-$^1$N-HSQC spectra of MAK33 V_{\text{L,\kappa}}-WT and MAK33 V_{\text{L,\kappa}}-I2E. Both spectra were acquired at a protein concentration of 50 $\mu$M in 20 mM phosphate, 50 mM NaCl (pH 6.5) at 298 K on a 600 MHz spectrometer equipped with a cryoprobe. b, chemical shift changes of MAK33 V_{\text{L,\kappa}} caused by the I2E substitution. $^1$H and $^1$N backbone chemical shifts were determined at a protein concentration of 50 $\mu$M at 298 K in 20 mM phosphate, 50 mM NaCl (pH 6.5). c, effects of I2E substitution in MAK33 V_{\text{L,\kappa}}. Strongly affected residues are marked on the MAK33 V_{\text{L,\kappa}} crystal structure (PDB code 1FHS). Ile-2 is shown in red, and residues with chemical shift changes $>0.05$ ppm are shown in dark purple, and unassigned residues are shown in gray.
Role of Residue 2 for Integrity of Antibody Variable Domains

and Ser-67 to Leu-73. The first region contains the structurally important residues Cys-23, which forms a conserved disulfide bond with Cys-88. The adjacent Arg-24 is involved in a conserved salt bridge with Asp-70 and the conserved Trp-35 lies in the core of Ig domains (50). Furthermore, the I2E substitution causes structural changes also at distant residues like Tyr-86 and Phe-87 (Fig. 8c). In consequence, a rearrangement of the hydrophobic core involving residues Ala-25, Ile-29, Trp-35, Leu-73, and Tyr-86, which exhibit major chemical shift changes, is induced. The CDRs also show shift perturbations, especially the loop including residues Arg-24 to Ile-29, which is close to the N terminus.

To gain further insight in the dynamics of the $V_{\kappa}$ domains, MD simulations in explicit solvent were performed on the MAK33 $V_{\kappa}$-WT and 1OPG $V_{\kappa}$-WT domains and on the MAK33 $V_{\kappa}$-I2E and 1OPG $V_{\kappa}$-E2I mutants. The substitutions were generated in silico based on the native crystal structures followed by extensive energy minimization before starting MD simulations (see “Experimental Procedures” for details). For all four cases, the r.m.s.d. of the backbone stayed close to the starting structure within $<2.0$ Å (Fig. 9). In the variants with an Ile at position 2 (MAK33 $V_{\kappa}$-I2E and 1OPG $V_{\kappa}$-WT), significant deviations and fluctuations of the Glu-2 side chains with respect to the starting structure were observed (red curves in Fig. 10, a and b). Deviations of up to 6–8 Å from the initial placement were observed that correspond to a partial or full dissociation of the Glu-2 side chain from the cavity, thereby adopting a fully solvent-exposed conformation (compare Fig. 10, d and e). The frequent disruption of contacts between Glu-2 and other side chains indicates that Glu does not contribute significantly to the stability.

FIGURE 9. a, comparison of backbone root mean square deviations (RMSD) from the corresponding experimental start structure versus data gathering simulation time for MAK33 $V_{\kappa}$-WT (black curve) and MAK33 $V_{\kappa}$-I2E (red curve). b, same for 1OPG $V_{\kappa}$-WT (red curve) and 1OPG $V_{\kappa}$-E2I (black curve). c, comparison of root mean square fluctuations of amino acid residues along the domain chain for 200-ns simulations of MAK33 $V_{\kappa}$-WT (black curve) and MAK33 $V_{\kappa}$-I2E (red curve). d, same for 1OPG $V_{\kappa}$-WT (red curve) and 1OPG $V_{\kappa}$-E2I (black curve).

The analysis of root mean square fluctuations of each residue with respect to the mean structure (Fig. 9, c and d) showed similar fluctuations for wild type and corresponding mutant chains. However, significantly larger fluctuations were observed for variants with a Glu at position 2 compared with those with Ile-2. Slightly increased fluctuations were also seen at loop segments that flank residue 2 (e.g. the loop around Ala-25), also identified in NMR experiments to have strong chemical shift perturbations. To quantify the relative stability of solvent-exposed and buried states of the side chain at residue 2, US simulations were performed to induce a dissociation of the side chain from the N-terminal cavity region. The distance between the Cα atom of residue 2 and the Cα atom of a residue at the floor of the N-terminal contact region served as reaction coordinate (see “Experimental Procedures” for details). The US simulations allowed for the calculation of a potential-of-mean force or associated free energy change for the dissociation of residue 2 (Fig. 10c). A distance of $\sim 12$ Å corresponded to a location of residue 2 in contact with the residues forming the N-terminal cavity region (smaller distances resulting in steric repulsion). At distances of 16–18 Å, the side chain of residue 2
had lost contacts with the N-terminal cavity region (larger distances resulting in disruption of additional contacts of subsequent residues along the chain). The US simulations indicate a significantly higher dissociation free energy barrier for the case of an Ile at position 2 compared with Glu-2 in both MAK33 and 1OPG V_{\kappa}-WT domains. Because in the folded form the Ile-2 residue stays in an associated conformation, one can assign $-2.5$ kcal/mol ($\sim 10$ kJ mol$^{-1}$) as a free energy difference of dissociation compared with Glu-2 as a folding stabilization free energy contribution of the Ile-2 variants. This is in excellent agreement with our experimental values.

In addition to the $V_{\kappa}$ domains, the $V_{\lambda}$ and $V_{\mu}$ domains (taken from the x-ray structures of 1AQK (49) and 1VGE (39), respectively), were also analyzed. MD simulations were performed starting from the x-ray conformations of the wild type structures and from variants with a Glu at position 2. In all simulations, the domain conformations stayed close (within $\sim 2$ Å) to the start structure for a 50-ns sampling time (Fig. 11, a and b). In the case of 1AQK $V_{\lambda}$, the Asn-2 wild type residue is partially solvent-exposed forming contacts with polar and charged residues near the N terminus (Fig. 10h). During the simulations, it transiently dissociates to more solvent-exposed states. Similarly, the Glu-2 variant (1AQK $V_{\lambda}$-N2E) also shows fluctuations to partially solvent-exposed states with limited contacts to the N-terminal region of the protein (Fig. 11c). In the case of 1VGE $V_{\mu}$, residue 2 is in a fully solvent-exposed state.
already in the x-ray starting structure (Fig. 10g). No cavity near the N terminus is present that could serve as a stable anchor region for residue 2 as found for the /H9260 domains (Fig. 10f). Both the simulations of 1VGE VH-WT and the V2E mutant showed large fluctuations of residue 2 and no stable binding mode could be identified (Fig. 11d). Thus, in these cases neither the wild type nor the Glu-2 variants adopt a stable state with a buried side chain of residue 2 that could contribute to the stability of the protein, in contrast to what was observed for the Ile-2 variants of MAK33 and 1OPG VL /H9261. This is largely due to the absence of an appropriate cavity region near the N terminus and explains why mutation of residue 2 has little or no influence on protein stability in the case of the VL /H9261 and VH domains.

**DISCUSSION**

We show here that residue 2 is crucial for the integrity and stability of VL domains of the /H9260 family, and this residue is a key factor that controls its amyloidogenic properties. Specifically, uncharged residues at position 2 of VL /H9260 domains are important for the maintenance of the structural integrity and a high stability. In agreement with their extremely low frequency in naturally occurring VL /H9260 domains, charged residues do not support this.

During NMR backbone assignment, we encountered several regions that are exchanged-broadened and therefore could not be assigned completely. Interestingly, these regions match well with dynamic residues in the dimer interface of the human VL /K sequence LEN, as identified by Mukherjee et al. (51). In this VL domain (LEN), residues 1–9, 37–58, and 89–100 were shown to exhibit pH-dependent millisecond dynamics, which seem to be related to the process of amyloid fibril formation. The presence of residue 2 within these dynamic regions is another indicator for its relevance for amyloidogenicity. This is supported by our MD simulations that show an increase in N-terminal dynamics caused by a charged residue at position 2. Overall, the I2E substitution affects large parts of the VL /K structure. Both the disruption of the conserved salt bridge and the changes in the hydrophobic core presumably contribute to the loss of thermodynamic stability. However, the short strand-connecting helix (residues 79–83), which is highly conserved in antibody variable domains and known to be important for early folding events of the domain (13) was not affected by the I2E substitution, implying that this replacement may not be involved in the amyloidogenicity of the VL variants studied here.

Hydrophobic core packing is well known to be an important stabilizing factor in proteins in general (52–54) and antibody domains in particular (55). Therefore, the destabilizing effects of charged residues at position 2 could be explained by the fact that they are energetically unfavorable within the hydrophobic core. A charged residue at position 2 of a VL /K domain leads to a partial exposure of the hydrophobic interior and to the disruption of an optimal packing around residue 2. This in turn causes rearrangements of the whole hydrophobic core region. Consequently, this results in the destabilization of the fold and an increased aggregation propensity. For the VL / and VH domains, the residue at position 2 is largely solvent-exposed or forms only transient polar contacts with the surrounding protein surface. In MD simulations, both the wild type residues at position 2 or substitutions with Glu resulted in fluctuating solvent-exposed conformations due to the lack of a stable binding region. Hence, in the case of the VL / and VH domains, residue 2 is not part of the hydrophobic core, and substitution of this residue also does not disrupt the hydrophobic packing of the
domains. Whereas the binding of the N-terminal segment to the hydrophobic cavity is crucial for stability in $V_{L,K}$ domains, it appears to be less important for $V_{L,A}$ and $V_{L,H}$ domains. Unlike in the $V_{L,K}$ domains where the hydrophobic cavity requires binding of the N-terminal residue 2 to prevent the unfavorable exposure of nonpolar residues, in the $V_{L,A}$ and $V_{L,H}$ domains, the respective hydrophobic cavities are less hydrophobic and do not require binding of residue 2 for protection.

It has been suggested that the formation of amyloid fibrils does not begin from the native state of a protein but more likely from a partially folded state or an intermediate state (32, 56, 57). In this context, the occurrence of particularly nonconservative mutations that involve a change in side chain chemistry in a structurally important region and the correlation with an increased propensity to form amyloid have been reported (32, 58). Mutations and changes in conditions that lead to destabilization of the native state might therefore serve to increase the population of an intermediate state or partially folded species, thereby enabling more molecules to be diverted into an amyloid-forming pathway (59, 60). In this study, the less stable MAK33 and 10PG $V_{L,K}$ variants readily formed fibrils, whereas their stable counterparts did not. Although stability is an important factor determining the amyloidogenic propensity of proteins, the amino acid sequence plays a major role (61) as this determines the amyloid fibril formation kinetics and the types of intermediates populated (13, 32, 56). Although some residues protect against amyloid formation, their chemistry and position within the protein appear to be more important. This protective role is largely attributed to edge strands of $\beta$-sheet proteins (62–64). Residue 2 in antibody variable domains appears in $\beta$-strand A, which is one of four edge strands (A, D and C*, G) of the $\beta$-sandwich topology. Uncharged residues at this position might have evolved to prevent undesirable $\beta$-sheet self-propagation (63). The frequent dissociation and solvent exposure of charged residues at position 2 of $V_{L,K}$ domains presumably lead to the disruption of native H-bonds and the subsequent exposure of H-bond donors and acceptors on the edge strand interface. This might result in $V_{L}$ edge-to-edge aggregation (65). The rare occurrence of charged residues at this position within edge strand A and the high amyloidogenic propensity of these $V_{L,K}$ variants are in line with this explanation. Although there exists a huge similarity between the three different families of variable domains, it has also been shown that they differ significantly in their biophysical properties and propensity to aggregate to amyloid fibrils (55, 66). Our study identifies an important property of the $V_{L,K}$ family, not pertaining to the $V_{L,A}$ and $V_{L,H}$ domains, which controls its amyloidogenicity. A compelling amount of evidence from studies on sequence and structural features that can predispose to amyloid formation has often been limited to given $V_{L}$ subfamilies (e.g. $\kappa$1 or $\lambda$6) with hardly one feature pertaining to all or an entire variable domain family (67–69). Our results identify a key feature that can destabilize and as a result predispose an entire variable domain family to fibrillar aggregation. The importance of residue 2 for the $V_{L,K}$ family determined here adds to the understanding of antibody LC amyloidosis linking the destabilization of native interactions with amyloid formation. In the context of AL amyloidosis, and also in the broader context of amyloid diseases, however, it remains to be seen whether point mutations promoting fibril formation generally affect stability. It seems reasonable to assume that other mechanisms may also apply.

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A Residue-specific Shift in Stability and Amyloidogenicity of Antibody Variable Domains
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