Daratumumab for Immunoglobulin Light Chain Amyloidosis

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Immunoglobulin light chain (AL) amyloidosis is characterized by small, indolent plasma cell clones that uniformly demonstrate high expression of CD38 surface molecule and produce toxic, free light chains. The use of daratumumab, a human anti-CD38 monoclonal antibody, has shown important efficacy with minor toxicities in patients with relapsed or refractory AL amyloidosis. Early data from the phase III ANDROMEDA clinical trial, which evaluated the combination of subcutaneous daratumumab with standard-of-care cyclophosphamide, bortezomib and dexamethasone in patients with newly diagnosed AL amyloidosis, showed an outstanding rate of deep haematological response with significant improvement in organ function, results that led to the approval of daratumumab for the treatment of the disease.

A new era for the management of AL has arisen and daratumumab is the centrepiece. In the current article, we provide an overview of the mechanism of action of daratumumab and review the clinical evidence that supports its use in the management of AL amyloidosis.

Keywords
Light chain amyloidosis, monoclonal antibody, daratumumab

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Immunoglobulin light chain (AL) amyloidosis is the most common type of systemic amyloidosis, a spectrum of diseases induced by misfolded proteins that generate insoluble amyloid fibrils and deposits in various organs. The precursor protein in AL amyloidosis is a toxic, free light chain (FLC) secreted by an indolent plasma cell or B-cell clone in the bone marrow. The most affected organs include the kidney, heart, liver, peripheral and autonomous nervous systems, and soft tissues. Heart involvement drives down survival of patients, according to the Mayo risk stratification system, which is based on cardiac biomarkers (N-terminal of the prohormone brain natriuretic peptide and high-sensitivity cardiac troponin T) and revised by the incorporation of difference of serum free light chain (dFLC). Treatment of AL amyloidosis is aimed at eradication of the underlying clone. The backbone of therapy includes regimens adopted from the field of multiple myeloma (MM). These antical agents include alkylators (melphalan and cyclophosphamide), proteasome inhibitors (bortezomib, carfilzomib and ixazomib) and immunomodulatory drugs (lenalidomide and pomalidomide). Early recognition and deep haematological responses, defined as very good partial response (VGPR) or better, have been correlated with improved overall survival (OS) over recent decades, with 2-year OS of 60% and 4-year OS of 54%. However, the disease is still incurable and 30% of patients will still die within 6 months of diagnosis, while a significant number are prone to toxicities related to conventional treatments. The development of more effective combinations that will further improve haematological responses will lead to prolonged survival and organ function restoration. Recently, the results of the phase III, randomized, open-label, multicentre ANDROMEDA (AMY3001) trial have been published. The study evaluated the combination of subcutaneous (SC) daratumumab with standard-of-care cyclophosphamide, bortezomib and dexamethasone (CyBoD) in newly diagnosed AL amyloidosis. Based on ANDROMEDA, the US Food and Drug Administration (FDA) and European Medicines Agency approved the use of SC daratumumab in combination with CyBoD for the treatment of newly diagnosed AL amyloidosis with Mayo stage I–III A, making it the first approved therapeutic regimen for this unique disease.

This article provides an overview of the mechanism of action of daratumumab, reviews the clinical evidence that supports its use in AL amyloidosis and describes how daratumumab could be incorporated into the current treatment strategy.

CD38 in AL amyloidosis clone

The biology of the plasma cell clone is considered to play a significant role in the pathogenesis of AL amyloidosis. The bone marrow usually shows low-grade plasma cell clones with low proliferation index. In fact, the presence of bone marrow plasma cells >10% is associated with poor survival regardless of Mayo stage. Clonal plasma cells produce large amounts of amyloidogenic FLCs that increase oxidative stress and render plasma cells highly dependent on proteasome activity. Thus, amyloidotic cells are intrinsically sensitized to proteasome inhibitors and bortezomib has become the centrepiece in AL amyloidosis management.
reveal that AL amyloidosis clones have more features in common with monoclonal gammapathy of undetermined significance than with MM, while the most frequent chromosomal aberration is translocation (11;14), which is also associated with adverse outcomes in patients treated with bortezomib-based regimens.

Studies using flow cytometry and next-generation sequencing have demonstrated that most clonal plasma cells express CD38, while higher expression has been correlated with advanced cardiac involvement and adverse prognosis in AL amyloidosis. CD38 is a 45-kD, type 2 transmembrane molecule with dual function as both transmembrane signalling receptor and as enzyme. Interaction with its ligand, CD31, leads to cell survival and migration. Moreover, CD38 acts as an ectoenzyme by converting nicotinamide adenine dinucleotide to cyclic adenosine diphosphate-ribose, which regulates calcium release and lymphocyte proliferation. CD38 is present on both normal and malignant plasma cells, but also on regulatory B cells, regulatory T cells and natural killer cells. The rationale for targeting the CD38 molecule in myeloma and other plasma cell dyscrasias lies in the fact that CD38 is ubiquitously expressed on plasma cells and on its role in cell signalling.

**Daratumumab mechanism of action**

The small size and the indolent nature of the AL amyloidosis plasma cell clone make therapy with monoclonal antibodies feasible and optimal. Daratumumab is a human IgGκ monoclonal antibody (MoAb) that binds with high affinity to CD38 on clonal plasma cells and other lymphoid cells. Similarly to antibodies, MoAbs have two distinct functional domains: a constant fragment (Fc) that connects to effector cells via Fcγ receptors (FcγR) and a variable fragment that binds to antigens. Daratumumab mediates cytotoxicity of plasma cells through pleiotropic mechanisms, including antibody-dependent cellular cytotoxicity (ADCC), antibody-dependent cellular phagocytosis (ADCP), complement-dependent cytotoxicity and direct apoptosis. Binding of the Fc domain to the CD38 epitope on clonal plasma cells promotes the activation of natural killer cells, which results in degranulation of cytotoxic cytokines and lysis of tumour cells by ADCC, and the activation of other effector cells (macrophages and monocytes), which leads to phagocytosis (ADCP).

Similarly, the interaction of daratumumab with C1q activates the classical complement pathway and the formation of membrane attack complex. Daratumumab induces programmed cell death of CD38+ myeloma cells through FcγR-mediated cross-linking; CD38 redistribution on the surface, loss of membrane integrity and morphological modifications contribute to programmed cell death activation. Daratumumab has immunomodulatory effects, too. The absolute number of natural killer cells is significantly reduced during daratumumab therapy. Regulatory T cells and other myeloid-derived suppressor cells are also depleted. At the same time, a substantial increase in CD4+ Th1 cells and CD8+ effector T cells could enhance an adaptive immune response. Finally, changes in enzymatic activity of daratumumab have also been studied. Inhibition of CD38 ectoenzyme function decreases the release of adenosine, an immunosuppressor that eliminates the activity of immune effector cells.

**Daratumumab in AL amyloidosis**

Daratumumab monotherapy has produced durable responses with a manageable toxicity profile in relapsed/refractory MM. The addition of daratumumab to lenalidomide or bortezomib significantly prolonged progression-free survival (PFS) and improved the overall response rate of patients who had received at least one prior treatment. Among patients with newly diagnosed MM, both eligible and ineligible for autologous stem cell transplantation (ASCT), daratumumab in combination with standard-of-care regimens improved OS and PFS. Furthermore, patients achieved deeper responses, and evaluation of minimal residual disease (MRD) showed higher MRD-negative rates.

The results of the COLUMBA study showed that SC daratumumab (1,800 mg in 15 mL) with recombinant human hyaluronidase PH20, was non-inferior to intravenous (IV) daratumumab in terms of efficacy and pharmacokinetics, and led to FDA approval. The most common side effects of daratumumab are infusion-related reactions and infections.

In AL amyloidosis, daratumumab was initially introduced in the relapsed setting. The first report was published in 2016. Two patients with previously treated AL amyloidosis received daratumumab with dose and schedule similar to those used for myeloma approval. Both patients achieved haematological complete remission soon after they began treatment. One of the patients had cardiac involvement and tolerated the IV infusion of daratumumab, suggesting that the treatment may also be an option for frail patients. A retrospective analysis of 25 heavily pretreated patients, published in 2017 by Kaufman et al. showed that daratumumab produced rapid and deep responses. The overall haematological response rate was 76% (complete response [CR] in 36% and VGPR in 24%), and median time to response was 1 month. Responses were actually more favourable than those with other available regimens in the relapse setting, and adverse events were similar to those observed in MM. Further small cohorts of patients with relapsed AL amyloidosis who received IV daratumumab had extraordinary haematological responses (63–100%), time to response (1–4 weeks) and organ response (cardiac response in up to 55% and renal response in up to 67%). A large cohort of patients (n=168) compared daratumumab monotherapy with the combination of daratumumab plus bortezomib and dexamethasone. Results were promising, with 64% versus 66% of patients achieving CR, showing that combination therapies are feasible in relapsed AL amyloidosis. Daratumumab has been an effective option even for patients with high tumour burden or nephrotic-range proteinuria. Moreover, daratumumab in combination with immunomodulatory drugs appears to be safe and more effective than in MM patients. These results provided the rationale for the ANDROMEDA study in newly diagnosed patients.

**Daratumumab in combination with cyclophosphamide, bortezomib and dexamethasone in newly diagnosed AL amyloidosis**

In the first-line setting, IV daratumumab was first administered in two patients with advanced cardiac involvement and they both achieved complete remission without serious adverse events. The phase III ANDROMEDA study demonstrated that the addition of SC daratumumab to standard CyBorD improved haematological and organ responses. Patients were randomized to receive either CyBorD alone for six cycles, or CyBorD plus daratumumab (DARA-CyBorD) for six cycles followed by maintenance therapy with monthly daratumumab for up to 24 cycles. Eligible patients had to have histopathological diagnosis of AL amyloidosis (based on immunohistochemistry or electron microscopy) with measurable disease and at least one involved organ according to International Society of Amyloidosis (ISA) criteria. Adequate organ function (estimated glomerular filtration rate of at least 20 mL/ min/1.73 m²) was required, and patients were not eligible if they were Mayo stage IIIb (NT-proBNP level >8,500 ng/L) or if they had other severe cardiovascular diseases.

The primary endpoint of the study was overall haematological CR rate. As a human IgGκ MoAb, daratumumab is detected by serum immunofixation electrophoresis, making CR assessment according to ISA
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In the first interim analysis of the ANDROMEDA study, the median follow-up was 11.4 months. Overall, 388 patients were randomized (195 to the DARAYCyBorD group and 193 to the CyBorD-only control group), and haematological CR rate was 53.3% in the DARAYCyBorD group versus 18.1% in the CyBorD group (relative risk ratio 0.49, 95% confidence interval [CI] 0.21–1.1; p = 0.004). At the 6-month landmark analysis, haematological CR rate was 49.7% in the DARAYCyBorD group versus 14% in the CyBorD group. In the DARAYCyBorD group, median time to first response (at least partial response) was 9 months and median time to VGPR was 19 days. Importantly, the median time to haematological CR was 60 days in the DARAYCyBorD group versus 85 days in the CyBorD group. The rate of VGPR or better was 78.5% and 49.2% respectively (p = 0.0003), while deep haematological responses with dFLC ≤20 mg/L or iFLC ≤20 mg/L occurred more frequently with the daratumumab combination treatment (70.8% versus 20.2% and 64.1% versus 30.6%, respectively). Independently of the criteria used, there was a substantial benefit in terms of deep haematological responses with the addition of daratumumab. In an updated analysis that was presented later, the median follow-up was 20.3 months, and haematological CR rates were significantly higher in the DARAYCyBorD group (59% versus 19%; odds ratio 5.9, 95% CI 3.7–9.4; p < 0.0001).

Improvement of organ function is also a major goal of AL amyloidosis therapy. Patients who attain deep haematological responses are more likely to obtain an organ response and this can substantially improve patients’ quality of life. Patients with measurable disease who achieve CR or very low FLC levels and patients without measurable disease after the completion of therapy should be evaluated for MRD using next-generation flow cytometry or next-generation sequencing. Although the MRD data from the ANDROMEDA trial are not yet available, MRD-negative status in other series has been associated with very low probability of haematological relapse and with high rates of organ responses. These results highlight that MRD negativity should be the optimal goal of AL amyloidosis anticlonal therapy. Furthermore, as next-generation flow methods are increasingly available, a shift towards more individualized treatments is anticipated. Other antigens on the surface of plasma cells that have been targeted initially in myeloma but are also tested in AL amyloidosis, include CD20 surface molecule (targeted by rituximab), Signalling Lymphocyte Activation Marker Family member 7 (SLAMF7) receptor (targeted by elotuzumab) and B-cell maturation antigen (targeted by belantamab mafodotin antibody-drug conjugate).

Future perspectives

The combination of bortezomib with cyclophosphamide and dexamethasone (CyBorD) results in VGPR/CR rates of 43%, with organ responses in up to 25% of patients, and was considered the mainstay of AL amyloidosis therapy until recently. The ANDROMEDA trial demonstrated that the addition of daratumumab to CyBorD, yields faster, deeper and more durable responses with limited toxicity. As a result, daratumumab has moved to the frontline of AL amyloidosis therapy and DARAYCyBorD has become the new standard of care.

A very appealing feature of daratumumab is its significant ability to induce deep haematological responses within the first month of its administration. Findings from the ANDROMEDA trial along with previously published data, suggest that the achievement of stringent dFLC or iFLC responses, even without fulfilling ISA criteria for CR, is associated with significantly improved survival and higher organ responses. As mentioned above, as a human IgGκ mAb, daratumumab is detected by serum immunofixation electrophoresis, making CR assessment challenging. Patients with measurable disease who achieve CR or very low FLCs levels and patients without measurable disease after the completion of therapy should be evaluated for MRD using next-generation flow cytometry or next-generation sequencing. Although the MRD data from the ANDROMEDA trial are not yet available, MRD-negative status in other series has been associated with very low probability of haematological relapse and with high rates of organ responses. These results highlight that MRD negativity should be the optimal goal of AL amyloidosis anticlonal therapy.
In AL amyloidosis, translocation (11;14) is the most common chromosomal alteration among patients (up to 50%). Carrying t(11;14) is an adverse prognostic factor, with inferior haematological responses in patients treated with bortezomib-based regimens (VGPR or better 52% versus 77%; p=0.004) and with shorter survival (median OS 15 versus 27 months; p=0.05). In the ANDROMEDA trial, haematological CR rates were independent of the presence of t(11;14) (55% versus 52%) in the DARA-CyBorD group. In fact, the addition of daratumumab may overcome the adverse impact of t(11;14), rendering daratumumab a promising tool for this subgroup of patients, either as monotherapy or in combinations. In addition, t(11;14) is associated with overexpression of anti-apoptotic factor Bcl-2, which is targeted by the Bcl-2 inhibitor, venetoclax. However, based on data from myeloma patients, high mortality rates raise concerns about safety in this frail population; some case reports have implied that the addition of venetoclax to daratumumab could enhance the outcome in patients with relapsed/refractory AL amyloidosis.

A very high-risk cohort of patients includes Mayo stage IIIb (NT-proBNP levels >8500 pg/mL) and patients with extremely low systolic blood pressure (<100 mmHg), with median survival of 3–6 months. The early eradication of the toxic FLCs and the underlying plasma cell clone is crucial. However, standard treatment with CyBorD is ill-tolerated. Daratumumab, on the other hand, is a regimen with manageable adverse events, no cardiotoxicity, and the introduction of SC administration significantly reduces the risk of volume overload. Given the high rates of substantial deep responses and the high rates of cardiac responses in relapsed AL but also in the ANDROMEDA study, daratumumab has emerged as an attractive option for high-risk patients. An ongoing phase II trial is currently assessing the safety and efficacy of daratumumab monotherapy in previously treatment-naive stage IIIb AL amyloidosis patients.

The rapid reduction of the toxic FLCs is also relevant for the restoration of kidney function in patients with renal amyloidosis. The renal damage often leads to end-stage renal disease; therefore, the use of daratumumab, which reduces CRs and VGPRs in a short time, in combination with bortezomib, which has a protective effect on the kidney, offers a very promising novel option. Daratumumab-based regimens have also been considered for the treatment of the rare entities of monoclonal gammapathies of renal significance, with great efficacy and safety in a small cohort of 25 patients. Although monoclonal antibodies are safe for the treatment of patients with renal dysfunction, in a series of 168 patients with AL amyloidosis treated with daratumumab-based regimens, nephrotic-range proteinuria was an adverse factor for haematological responses.

Besides cardiac and renal amyloidosis, there is also concern for the management of patients with mild-to-severe peripheral nerve involvement. Owing to the neurotoxicity of bortezomib and fears of late-onset peripheral neuropathy, daratumumab could be as high as 20%, and the introduction of novel immunotherapies makes its role debatable. In fact, the attainment of deep haematological and organ responses with the use of daratumumab-based regimens as a first-line agent (CR 53%, cardiac response 42% and renal response 53%), together with its favourable safety profile, eliminate the need for ASCT in AL amyloidosis, even for transplant-eligible patients. Depending on the depth of haematological response and the achievement of organ response post-daratumumab, ASCT could be considered for those patients with residual disease or persistent organ dysfunction. Longer follow-up of the ANDROMEDA study will further elucidate the role of daratumumab maintenance therapy as a more appropriate chemotherapy-free strategy, compared with ASCT, for patients who have not achieved complete eradication of clonal cells.

In the setting of relapse, the establishment of DARA-CyBorD as the mainstay of first-line treatment raises new questions about the management of relapsed or refractory AL amyloidosis patients. Today, many patients are still naïve to daratumumab. Thus, single-agent therapy or combination with low-dose lenalidomide should be considered. Overall, large prospective clinical trials with modern regimens and novel combinations are needed to address these new challenges. The management strategy should always be tailored according to risk stratification systems, organ involvement, clonal cell biology and haematological response.

Conclusions

The overexpression of CD38 surface molecule has provided the rationale for using daratumumab in the treatment of patients with AL amyloidosis. The results of the ANDROMEDA study show that deep and durable haematological and organ responses are feasible with the addition of daratumumab to CyBorD and this led to the first approved treatment for the disease. Early diagnosis and immediate initiation of therapy are still the mainstay of AL amyloidosis management. The high efficacy and tolerability of daratumumab, even in high-risk populations, highlight a promising new era for the treatment of AL amyloidosis.

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