

Adult-onset autoimmune diabetes: current knowledge and implications for management

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Abstract | Adult-onset autoimmune diabetes is a heterogeneous disease that is characterized by a reduced genetic load, a less intensive autoimmune process and a mild metabolic decompensation at onset compared with young-onset type 1 diabetes mellitus (T1DM). The majority of patients with adult-onset autoimmune diabetes do not require insulin treatment for at least 6 months after diagnosis. Such patients are defined as having latent autoimmune diabetes in adults (LADA), which is distinct from classic adult-onset T1DM. The extensive heterogeneity of adult-onset autoimmune diabetes is apparent beyond the distinction between classic adult-onset T1DM and LADA. LADA is characterized by genetic, phenotypic and humoral heterogeneity, encompassing different degrees of insulin resistance and autoimmunity; this heterogeneity is probably a result of different pathological mechanisms, which have implications for treatment. The existence of heterogeneous phenotypes in LADA makes it difficult to establish an *a priori* treatment algorithm, and therefore, a personalized medicine approach is required. In this Review, we discuss the current understanding and gaps in knowledge regarding the pathophysiology and clinical features of adult-onset autoimmune diabetes and highlight the similarities and differences with classic T1DM and type 2 diabetes mellitus.

Autoimmune diabetes is a heterogeneous disease that is generally regarded as a condition that presents in childhood or adolescence. However, a substantial proportion of patients experience onset in adulthood¹. In this case, termed adult-onset autoimmune diabetes, the disease is even more heterogeneous than young-onset autoimmune diabetes, as the rate of β -cell destruction is highly variable, which is probably due to differences in the penetrance of genetic and immune factors^{2–4}. In the past few decades, particular attention has been paid to the characterization of adult-onset autoimmune diabetes. Epidemiological studies have highlighted that the majority of patients with onset in adulthood do not require treatment with insulin at the time of diagnosis, and these patients are defined as having latent autoimmune diabetes in adults (LADA)^{5–7}. This term was introduced in the early 1990s to define a subgroup of patients who had non-insulin-requiring diabetes mellitus that was initially thought to be type 2 diabetes mellitus (T2DM) but who had detectable serum immune markers of type 1 diabetes mellitus (T1DM)^{5,8}. Alternative eponyms for this condition have been proposed, such as ‘type 1.5 diabetes’ (REF. 9), ‘non-insulin-requiring autoimmune diabetes’ (REF. 10) and ‘slowly progressive insulin-dependent type 1 diabetes’ (REF. 11).

In 2005, the Immunology of Diabetes Society proposed three main criteria for the diagnosis of LADA: adult age of onset (>30 years); the presence of any islet autoantibody; and the absence of insulin requirement for at least 6 months after diagnosis¹². However, controversies regarding these criteria still exist. The major criticism is the subjectivity of the clinician’s decision to start insulin treatment, which is a key factor affecting the classification¹³. The American Diabetes Association and WHO do not recognize LADA as a distinct entity, as they include LADA within the T1DM classification^{14,15}. Owing to the slow rate of β -cell loss that is observed in LADA⁴, the period of insulin independence after onset can distinguish patients with LADA from those with classic adult-onset T1DM, who require insulin within 3 months of diagnosis¹⁴. The broad definition of LADA¹² — which includes any patient with diabetes who does not require insulin and who is positive for any islet autoantibody, regardless of titre, number or epitope specificity — represents the basic prerequisite for the wide degree of heterogeneity observed in this form of diabetes.

New data that further elucidate the pathophysiology and clinical implications of adult-onset autoimmune diabetes have been reported. Nonetheless, these topics

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Key points

- Adult-onset autoimmune diabetes encompasses a wide spectrum of heterogeneous genotypes and phenotypes, ranging from classic adult-onset type 1 diabetes mellitus to latent autoimmune diabetes in adults (LADA)
- The heterogeneity of LADA arises from its definition as being present in any adult with diabetes who does not require insulin and who is positive for any islet autoantibody, regardless of titre, number or epitope specificity
- The heterogeneity of LADA manifests in different clinical phenotypes, ranging from prevalent insulin resistance to prevalent insulin deficiency, each of which might be associated with different autoimmune and metabolic markers
- Although patients with LADA are leaner and have healthier lipid and blood pressure profiles, evidence shows that there is no difference in cardiovascular outcomes between these patients and those with type 2 diabetes mellitus
- The extensive heterogeneity of adult-onset autoimmune diabetes, and particularly LADA, makes it difficult to determine an *a priori* algorithm for treatment
- The successful treatment of adult-onset autoimmune diabetes will require a personalized medicine approach that takes into account the intrinsic characteristics of each patient

are still under debate, and controversies in the classification of adult-onset autoimmune diabetes remain. In this Review, we report and critically discuss the results of the most recent and relevant studies, emphasizing the differences between T2DM, LADA, young-onset T1DM and adult-onset T1DM. Data regarding the epidemiology of adult-onset autoimmune diabetes are gathered to define the global picture of this disease. We review the pathogenesis of this disease in depth, highlighting genetic susceptibility and immune features as well as their influence on β -cell function. In addition, the metabolic and clinical features of individuals with LADA from different ethnic groups are summarized, and the data regarding the risk of both microvascular and macrovascular complications are discussed. Finally, we review the most recent advances towards a pathophysiology-oriented treatment paradigm for LADA.

Epidemiology

The growing body of literature on the epidemiology of adult-onset autoimmune diabetes is a testament to the increasing interest in adult-onset autoimmune diabetes. The available data show that adult-onset T1DM is more common than previously recognized. Indeed, approximately 40% of T1DM cases globally occur in people older than 30 years of age¹. In Italy, the incidence of T1DM in individuals aged 30–49 years is similar to that in adolescents aged 15–19 years¹⁶. Moreover, a study carried out in Sweden showed that the real incidence of T1DM in individuals aged 15–34 years was twofold to threefold higher than previously reported¹⁷. Two large studies published in the past 5 years have investigated the relative frequencies of T1DM autoantibody positivity in classic adult-onset T1DM and LADA and reported 3.3-fold and 12.8-fold higher frequencies of LADA in people of European and Arab origin, respectively^{6,7}. The results of both studies suggest that LADA is the most frequent form of adult-onset autoimmune diabetes. In addition, multicentre studies have reported that 4–14%

of patients with a diagnosis of T2DM are positive for T1DM-associated autoantibodies, which is indicative of a diagnosis of LADA^{6,7,18–27}.

The frequency of patients with T1DM-related autoantibodies among all patients diagnosed with T2DM varies considerably depending on ethnicity (TABLE 1), with the highest rates of LADA reported in people of northern European origins (7–14%)^{18,19,25,28}. The NonInsulin Requiring Autoimmune Diabetes (NIRAD) Study²², carried out in Italy, found that the cumulative frequency of positivity for either glutamic acid decarboxylase (GAD) autoantibodies and/or tyrosine phosphatase IA-2 (IA-2) autoantibodies was 4.5% in a cohort of 5,330 patients with T2DM²². Testing for zinc transporter 8 (ZnT8) autoantibodies identified an additional 1.4% of patients as autoantibody positive, bringing the potential frequency of autoantibody positivity in this T2DM cohort to 5.9%²⁹. A frequency of 9.7% was reported in Action LADA, a European multicentre study that evaluated 6,000 patients with adult-onset diabetes who attended primary and secondary care centres⁶. In the multicentre LADA China Study, the frequency of autoantibody positivity among adults with T2DM was reported to be 5.9%, which is similar to, or even higher than, the frequency described in European countries²⁶. This observation was quite unexpected, considering that childhood-onset T1DM is rare in China³⁰. Outside of Europe, the highest frequency of autoantibody positivity was reported in Indonesia, with up to 20% of T2DM patients being affected³¹, whereas the lowest frequencies were reported in Alaska³² and Papua New Guinea³³. Studies have also reported that African-American, Hispanic and Arab people have a lower prevalence of adult-onset autoimmune diabetes than white people^{7,34}.

The worldwide variance observed in the frequency of autoantibody positivity in patients with T2DM could be primarily due to differences in study design and inclusion criteria (such as age at diagnosis, sex, mode of recruitment, number and type of autoantibodies tested and sensitivity and specificity of autoantibody assays) as well as ethnicity. Furthermore, the increasing prevalence of T2DM in some populations could influence the frequency of autoantibody positivity in patients with T2DM³⁵.

Genes, autoimmunity and β cells

Adult-onset autoimmune diabetes has a lower 'genetic load' and is characterized by fewer diabetes-associated autoantibodies than young-onset T1DM^{2,3,36}. These genetic and autoimmune characteristics are consistent with a less severe functional deterioration of the β cells at disease onset than in young-onset T1DM³. Compared with young-onset T1DM, LADA represents the other extreme of the autoimmune diabetes spectrum, whereby genetic susceptibility, an autoimmune response and non-insulin-requiring presentation converge in a mild form of diabetes mellitus²² (TABLE 2).

Genetics

In LADA, genetic susceptibility has been investigated, but only in relation to genes that have been previously associated with young-onset T1DM^{2,37,38} or T2DM^{39,40}.

Table 1 | The prevalence of patients with T1DM-related autoantibodies among patients with T2DM

Study	Location	Type of study	Sample size (n)	Age (years)	Measured autoantibodies	Frequency of autoantibody positivity (%)	Refs
UKPDS 25	UK	Clinical-based	3,672	25–65	GAD and/or ICA	12	18
BOTNIA Study	Finland	Registry-based	1,122	28–83	GAD and/or IA-2	9.3	19
Eihme Study	Japan	Clinical-based	4,980	>20	GAD	3.8	20
ADOPT	USA, Europe	Clinical-based	4,357	30–75	GAD and/or IA-2	4.2	21
NIRAD Study	Italy	Clinical-based	5,330	30–75	GAD and/or IA-2	4.5	22
HUNT Study	Norway	Population-based	1,134	≥20	GAD	10	23
Tianjin	China	Population-based	8,109	≥15	GAD	9.2	24
Maioli <i>et al.</i>	Sardinia	Clinical-based	5,568	35–70	GAD	4.9	25
Action LADA	Europe	Clinical-based	6,810	30–70	GAD and/or IA-2, ZnT8	9.7	6
LADA China	China	Clinical-based	5,324	≥20	GAD	5.9	26
Maddaloni <i>et al.</i>	United Arab Emirates	Clinical-based	17,072	30–70	GAD and/or IA-2	2.6	7

ADOPT, A Diabetes Outcome Progression Trial; GAD, glutamic acid decarboxylase; HUNT, Nord-Trøndelag Health; IA-2, tyrosine phosphatase IA-2; ICA, islet-cell antibody; LADA, latent autoimmune diabetes in adults; NIRAD, Non Insulin Requiring Autoimmune Diabetes; UKPDS, United Kingdom Prospective Diabetes Study; ZnT8, zinc transporter 8.

This approach, which is based on the assumption that adult-onset autoimmune diabetes could have the same components of genetic susceptibility as young-onset T1DM, limits the possibility of discovering novel genes that are associated solely with adult-onset T1DM and/or LADA. In addition, most studies that have analysed genetic susceptibility did not compare patients with T1DM and LADA of the same age. Therefore, the data pertaining to this topic require careful interpretation.

The genes encoding human leukocyte antigen (*HLA*), cytotoxic T-lymphocyte antigen 4 (*CTLA4*), tyrosine-protein phosphatase non-receptor type 22 (*PTPN22*) and insulin (*INS*) have been associated with adult-onset autoimmune diabetes³. The *HLA-DRB1*04-DQB1*0302* and *HLA-DRB1*0301-DQB1*0201* haplotypes, which confer the highest susceptibility to T1DM^{41,42} and show a progressive decrease in frequency with increasing age at disease onset in children and adolescents^{43,44}, were shown to be further decreased in patients with older age at T1DM onset and were present at the lowest frequency in patients with LADA^{22,45}. The risk of T1DM that is conferred by the *CTLA4* Ala49Gly polymorphism in exon 1 (REF 37) did not decrease with age at clinical onset⁴⁶, suggesting that this polymorphism is also associated with LADA⁴⁷. The Cys185Thr single-nucleotide polymorphism in the *PTPN22* gene and the *INS* VNTR I/I genotype — which are both associated with increased susceptibility to T1DM^{38,48} — were present at a lower frequency in LADA than in young-onset T1DM^{45,49}. In addition, two different markers for the MHC class I polypeptide-related sequence A (*MICA*) gene, namely, the *MICA5* and *MICA5.1* alleles, have been shown to be associated with T1DM and LADA, respectively⁵⁰. The association between *MICA5.1* and LADA was later confirmed in a study carried out on patients from Finland⁵¹.

It is important to note that most of the studies were carried out in a single population. Furthermore, these studies were not well powered and thus did not have adequate statistical power to detect a true

association. Further studies with a larger number of patients are required to clarify the genetic associations that underpin LADA.

Autoimmunity

T1DM is a well-recognized cell-mediated autoimmune disease; in individuals with LADA, the presence of T cells that are reactive to islet-cell proteins provides some evidence of a cell-mediated immune response⁵². More specifically, the presence of insulinitis in LADA (as well as in T1DM) has been demonstrated by pancreatic scintigraphy with IL-2 radiolabelled with ^{99m}Tc, which revealed the presence of activated peripheral blood mononuclear cells in both T1DM and LADA⁵³. Importantly, it has been shown that a proportion of patients with phenotypic T2DM who are autoantibody negative display an autoimmune T-cell response, suggesting that these patients have autoantibodies that are currently undefined⁵⁴.

Islet autoantibodies are thought to be an epiphenomenon rather than key pathogenic factors in islet-cell destruction; however, they are used to discriminate autoimmune from non-autoimmune diabetes. The presence of GAD autoantibodies is not influenced by the age at disease onset and thus represents the most sensitive autoantibody marker in adult-onset T1DM and LADA¹⁸. Other autoantibodies are negatively associated with age at onset; older patients with T1DM are more likely than younger patients to be negative for insulin autoantibodies⁵⁵, IA-2 and ZnT8 (REFS 55,56), which are each detected only in small percentages of patients with LADA²⁹.

The islet autoantibodies that have been used thus far to identify patients with LADA are those that were first identified in T1DM. Autoantibodies against IA-2 are detected by different radioimmunoassays using different constructs (truncated fragments of the full-length protein that are used as antigens to detect autoantibodies) of the IA-2 protein⁵⁷. However, different autoantibodies against IA-2 exist, differing according to the protein

Table 2 | Genetic, metabolic and clinical features of LADA compared with T1DM and T2DM

Disease feature	Disease		
	T1DM	LADA	T2DM
Age at diagnosis	Childhood to adolescence and rarely in adulthood	>30 years	Adulthood and rarely in childhood to adolescence
Onset	Acute	Rarely acute	Slow
Autoimmunity	Severely increased	Increased	No change
Ketosis	Frequent	Rare	Rare
Insulin resistance	No change	Increased or no change	Severely increased
β -Cell function	Severely decreased	Decreased	Increased or no change
Insulin dependence	At onset	>6 months (even years) after onset	Years after onset
BMI	Underweight to normal	Normal to overweight	Overweight to obese
Risk of the metabolic syndrome	No change	Increased	Severely increased
HLA susceptibility	Severely increased	Increased	No change

LADA, latent autoimmune diabetes in adults; T1DM, type 1 diabetes mellitus; T2DM, type 2 diabetes mellitus.

epitope that they recognize. The NIRAD Study has shown that distinct constructs of the IA-2 protein might have different diagnostic sensitivities for the detection of IA-2 autoantibodies in T1DM and LADA⁵⁷. The intracytoplasmatic (_{IC}) IA-2_{IC(605–979)} construct showed the highest immunoreactivity in patients with T1DM, whereas the IA-2_(256–760) fragment showed the highest immunoreactivity in patients with LADA⁵⁷. This construct might represent a new sensitive diagnostic marker for the detection of islet autoimmunity in individuals with T2DM, which raises the possibility of the existence of different autoimmune processes that originate in individuals with obesity and insulin resistance⁵⁸. In addition, the Action LADA study found that the levels of pro-inflammatory cytokines such as IL-6 and tumour necrosis factor and the anti-inflammatory proteins IL-1 receptor antagonist and IL-10 were increased in patients with T2DM compared with individuals with autoimmune diabetes, whereas the levels of these cytokines were similar in LADA and T1DM⁵⁹.

β -Cell function

In young-onset autoimmune diabetes, β -cell function is already severely compromised at diagnosis⁶⁰. Early impairment of β -cell function is also seen in adult-onset autoimmune diabetes; however, this dysfunction is not as severe as in classic T1DM⁶¹. Indeed, a positive correlation between age at diagnosis of autoimmune diabetes and fasting C-peptide levels was reported⁶². Among adults diagnosed with autoimmune diabetes, those who fulfilled the diagnostic criteria for LADA had higher stimulated levels of C-peptide at all time-points following a mixed-meal tolerance test (MMTT)⁴. Prospective data from adults with diabetes who were followed up for 12 years from diagnosis showed that those with detectable autoantibodies had a severe degeneration of β -cell function, but this study did not distinguish individuals with LADA from those with adult-onset T1DM⁶¹. However, when

considering only individuals with LADA, C-peptide levels seemed to decline slowly⁴. Overall, residual C-peptide levels reflect the severity of disease progression, and progressive β -cell sparing is observed as the age of onset of autoimmune diabetes increases.

Metabolic and clinical heterogeneity

The intrinsic heterogeneity that is present in LADA arises from the original classification itself¹², which includes any patient with diabetes who is non-insulin requiring and who displays positivity for any islet autoantibody, regardless of titre, number or epitope specificity. Nevertheless, evidence exists showing that autoantibody titre and number are both related to different metabolic and clinical phenotypes of the disease, as outlined below. Moreover, new data suggest that the type of autoantibody that is present in patients with LADA indicates a different pathophysiology compared with that of classic T1DM.

Autoantibody titre and number

The presence of an insulin-deficient phenotype in patients with LADA who were in the highest tertile for GAD autoantibody levels was initially demonstrated in the BOTNIA Study¹⁹. Subsequently, the NIRAD Study²² highlighted the presence of a 'bimodal distribution' of the GAD autoantibody titre in patients with LADA — which had a nadir of 32 GAD autoantibody arbitrary units, corresponding to 300 WHO units — that identified two subpopulations, with high and low GAD autoantibody titres (FIG. 1). Based on the method used by Joanes and Gill⁶³, the bimodality of the GAD autoantibody titre in the NIRAD Study²² showed a bimodality coefficient ≥ 0.555 . The existence of subpopulations with a high or low GAD autoantibody titre has been confirmed by other, independent groups^{6,25,26}; however, some studies did not report a bimodal distribution of the GAD autoantibody titre¹⁹.

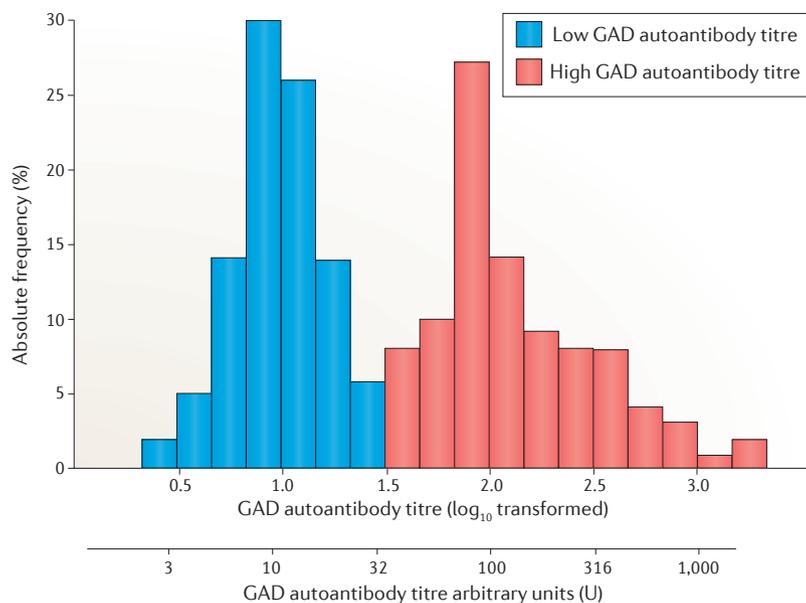


Figure 1 | Bimodal distribution of the glutamic acid decarboxylase autoantibody titre in patients with latent autoimmune diabetes in adults. The bar graph shows the frequency and distribution of the glutamic acid decarboxylase (GAD) autoantibody titre in patients with latent autoimmune diabetes in adults (LADA). The y axis shows the absolute frequency (%) of the values specified on the x axis. The x axis reports the log₁₀-transformed titre of GAD autoantibodies (and the respective titre in arbitrary units (U)) measured in 193 patients with LADA who participated in the Non Insulin Requiring Autoimmune Diabetes (NIRAD) Study²². The GAD autoantibody titre was measured using a radiobinding assay with *in vitro*-translated ³⁵S-methionine-labelled GAD65 antibody. The results were converted into U by extrapolation from a standard curve with a local standard designated 100 U. The graph illustrates the bimodal distribution of the GAD autoantibody titre in patients with LADA, with a nadir of 32 U, and identifies two subpopulations, with low (blue bars) and high (red bars) GAD autoantibody titres. The bimodality of the GAD autoantibody titre was verified using a bimodality coefficient⁶³, which we determined to be ≥ 0.555 . American Diabetes Association, Buzzetti, R. *et al.* High titer of autoantibodies to GAD identifies a specific phenotype of adult-onset autoimmune diabetes, *Diabetes Care*, 30, 932–938, American Diabetes Association, 2007. Copyright and all rights reserved. Material from this publication has been used with the permission of American Diabetes Association.

Compared with patients with LADA who had a low GAD autoantibody titre, those with a high titre had more severe autoimmunity, which resulted in higher levels of HbA_{1c}, a lower BMI and a lower prevalence of the metabolic syndrome²². In patients with LADA, these GAD-autoantibody titre-dependent differences in clinical and biochemical features were substantiated by genetic studies that reported that the frequencies of high-risk and moderate-risk *HLA* genotypes decreased linearly from a high to a low GAD autoantibody titre²². Similarly, the *PTPN22* risk genotype was also associated with a high GAD autoantibody titre in patients with LADA⁶⁴. Conversely, the transcription factor 7 like 2 (*TCF7L2*) risk allele for T2DM was associated with a low, rather than a high, GAD autoantibody titre⁶⁵. Of note, not all studies have confirmed the association between LADA and the *TCF7L2* gene⁶⁶; this discrepancy might be due to different patient inclusion criteria and the fact that such studies were underpowered. In addition, a higher frequency of autoantibodies associated

with autoimmune diabetes and other organ-specific autoantibodies — including thyroid peroxidase, steroid 21-hydroxylase, tissue transglutaminase and parietal cell autoantibodies — was detected in patients with LADA who had high GAD autoantibody titres than in those with low titres, thus confirming the increased severity of the autoimmune process in the former group of patients⁶⁷. These findings might have important clinical implications, and we suggest regular screening for other organ-specific autoantibodies (indicative of other autoimmune diseases) in patients with LADA according to the GAD autoantibody titre.

The number of islet autoantibodies in LADA has also been demonstrated to reflect the intensity of the autoimmune response and to predict future insulin deficiency^{18,68}. ZnT8 autoantibodies, as a marker in addition to GAD autoantibodies and IA-2 autoantibodies, have enabled stratification of the intensity of the islet autoimmune response, which is a clear reflection of the clinical phenotype of patients with adult-onset diabetes; features of more severe insulin insufficiency are proportional to the number of islet autoantibodies²⁹.

In addition, the United Kingdom Prospective Diabetes Study 25 reported that a high GAD autoantibody titre was associated with an increased risk of insulin requirement only among patients >55 years old at diagnosis¹⁸. However, conflicting results have since been reported; some studies demonstrated that a high titre of GAD autoantibodies was associated with a shorter insulin-free period^{19,69}, whereas other studies did not support this hypothesis^{25,70}. In the NIRAD Study, a high GAD autoantibody titre, BMI <25 kg/m², positivity for ZnT8 and IA-2_{IC(605–979)} autoantibodies and treatment with sulfonylurea in the first year after diagnosis led to markedly increased progression to insulin requirement in patients with LADA⁷¹.

Autoantibodies and pathophysiology

The IA-2_(256–760) autoantibody, which is present in 30% of patients with GAD autoantibody positivity and in 3.4% of patients who are negative for GAD and IA-2_{IC(605–979)} autoantibodies⁵⁷, was the only autoantibody that showed increased frequency with increasing BMI in a population of consecutive patients with T2DM⁵⁸. More interestingly, only patients with T2DM who were positive for the IA-2_(256–760) autoantibody showed clinical and metabolic phenotypes that exactly resembled those of patients with classic T2DM and obesity and had a substantially slower progression to insulin requirement within 7 years of follow-up than patients who were GAD autoantibody positive⁵⁸.

The presence of the IA-2_(256–760) autoantibody in patients with diabetes could underlie a pathophysiological mechanism resulting in a humoral immune response that is different from the response occurring in 'classic' autoimmune diabetes, as it is probably derived from the chronic systemic inflammation that is associated with obesity. Innate and acquired autoimmunity, specifically through the activity of macrophages and self-reactive T cells, contribute to the increased secretion of pro-inflammatory cytokines that are involved in

inflammatory processes⁷². The resulting inflammation might favour the presentation of extracellular antigens to antigen-presenting cells, thus promoting autoimmune activation⁷².

The IA-2 autoantibody in T1DM is directed against the intracellular portion of the protein⁷³. However, the epitope recognized by the IA-2₍₂₅₆₋₇₆₀₎ autoantibody is located in the extracellular domain of IA-2, which is more accessible to autoantibodies than intracellular epitopes. Thus, tissue damage induced by inflammation might trigger an autoimmune response to ‘cryptic’ self-antigens, thereby accelerating β-cell death. This hypothesis is supported by preliminary observations that the IA-2₍₂₅₆₋₇₆₀₎ autoantibody is detected in patients who are obese and do not have diabetes (R.B., unpublished observations). Therefore, as previously suggested⁴, the development of autoimmunity in patients with LADA could arise either as a consequence of the chronic inflammatory responses associated with obesity or as a result of a more specific environmental trigger that

can promote the activation of an autoimmune process similar to that which occurs in T1DM^{74,75} (FIG. 2). This hypothesis is purely speculative but encourages reflection on the existence of autoantibodies in T2DM and on the considerable heterogeneity of the patient population included in the definition of LADA.

The alternative possibility that the IA-2₍₂₅₆₋₇₆₀₎ autoantibody could reflect false-positive immunoreactivity was tested in a series of experiments that used different concentrations of unlabelled IA-2₍₂₅₆₋₇₆₀₎ fragments; the results demonstrated the specificity of antibody binding to ³⁵S-labelled IA-2₍₂₅₆₋₇₆₀₎ (REF. 57). Furthermore, a new autoantibody against the extracytoplasmic (EC) IA-2_{EC(26-577)} region can identify a novel antigenic determinant within the N terminus of the IA-2 protein⁷⁶. This autoantibody, whose epitope overlaps with the IA-2₍₂₅₆₋₇₆₀₎ epitope, can be detected in a subgroup of patients with adult-onset autoimmune diabetes who have a T2DM phenotype and who are negative for conventional islet autoantibodies⁷⁶. This finding adds further support to the relevance of the N-terminal region of IA-2 in terms of antigenicity in diabetes.

LADA versus T2DM

Although they are phenotypically distinct when large groups are compared, at the individual level, patients with LADA and patients with T2DM share clinical and metabolic characteristics, making it very difficult to diagnose LADA solely on the basis of the clinical phenotype. Therefore, individuals with LADA are often misdiagnosed as having T2DM. Several studies have been carried out to investigate the differences between LADA and T2DM with respect to genetic susceptibility and clinical features and have provided insight into how to correctly identify patients with LADA.

Genetic and clinical features

LADA shares several genetic features with T2DM. As previously mentioned, variants of the *TCF7L2* gene are also associated with LADA⁷⁷; this association is particularly strong in patients with a low GAD autoantibody titre⁶⁵. The *FTO*⁶⁶ gene is also associated with LADA, but to a lesser extent than *TCF7L2* (REF. 77). These associations suggest that LADA represents a genetic admixture of T1DM and T2DM.

Overall, patients with LADA show metabolic and clinical phenotypes that are substantially different from those of classic T2DM; relative to T2DM, LADA is characterized by higher fasting levels of glucose and HbA_{1c}, a lower prevalence of the metabolic syndrome, a higher frequency of thyroid peroxidase autoantibodies²², a higher frequency of *HLA* risk haplotypes and a consistently greater likelihood of insulin requirement^{6,71} (TABLE 2). In addition, patients with LADA had a lower stimulated C-peptide response at all time-points during an MMTT than individuals with T2DM, indicating reduced β-cell function⁴. This observation is in agreement with previous studies^{61,78}.

Although they are still present, these differences are less evident in patients with LADA who have a low GAD autoantibody titre or positivity for a single

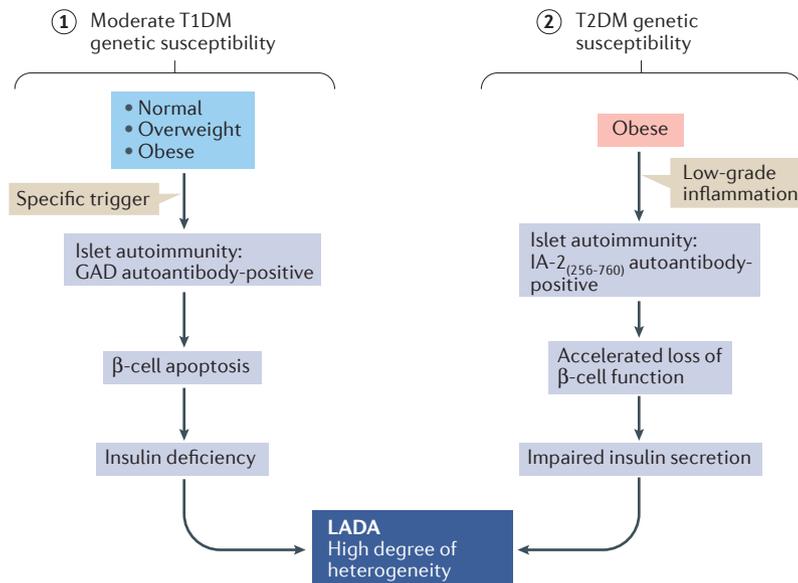


Figure 2 | Potential pathological mechanisms of latent autoimmune diabetes in adults. Here, we describe our working hypothesis regarding the pathophysiology of latent autoimmune diabetes in adults (LADA). In patients with moderate genetic susceptibility to type 1 diabetes mellitus (T1DM), specific immunological factors that are not well characterized can trigger an autoimmune process against the islets of Langerhans; this is independent of obesity, as it can occur in individuals with a normal BMI or who are overweight. This autoimmune process is marked by the appearance of glutamic acid decarboxylase (GAD) autoantibodies in the serum. Islet autoimmunity causes β-cell apoptosis, leading to insulin deficiency, which finally causes disease onset (1). LADA might also develop in individuals with obesity who have genetic susceptibility to type 2 diabetes mellitus (T2DM). The low-grade inflammation that characterizes visceral adiposity might trigger a low-grade autoimmune process that leads to the development of less severe islet autoimmunity, marked by the presence of serum autoantibodies against the tyrosine phosphatase IA-2₍₂₅₆₋₇₆₀₎ construct. This autoimmunity causes accelerated loss of β-cell function and impaired insulin secretion, which, when combined with the insulin resistance that commonly occurs in patients with obesity, causes hyperglycaemia and onset of diabetes mellitus (2). Our proposed pathological pathways explain the heterogeneous metabolic and clinical phenotypes of LADA, ranging from patients who are lean and insulin sensitive to those with obesity and insulin resistance, with a phenotype that is undistinguishable from that of T2DM.

islet-cell autoantibody (either GAD, IA-2_{IC} or ZnT8 autoantibodies)^{22,27}. Such data indicate that even a low GAD autoantibody titre or positivity for a single marker of autoimmunity can discriminate between autoimmune and non-autoimmune diabetes. Moreover, this observation suggests that autoimmunity is present in these patients, albeit at a low intensity. Furthermore, in the NIRAD Study²², serum samples from patients with GAD autoantibody positivity were tested in an inhibition assay, with an excess of unlabelled antigen in all samples; the results showed that in most cases, antibody binding was specific for GAD, lowering the possibility that the detection of low-intensity autoimmunity could be a false-positive result. In addition, the possibility of inclusion of some patients with T2DM in the LADA group in these studies cannot be ruled out, which is probably dependent on the sensitivity and specificity of the assays used.

Conversely, patients who are positive for only the IA-2₍₂₅₆₋₇₆₀₎ autoantibody do not differ from patients with T2DM with respect to clinical or metabolic phenotype⁵⁸. The only distinctive trait that can distinguish patients with positivity for the IA-2₍₂₅₆₋₇₆₀₎ autoantibody and T2DM is the frequency of IA-2_{IC(605-979)} autoantibody positivity⁵⁷; as discussed above, this finding implies the presence of autoimmunity, which is possibly the result of a different pathogenic mechanism between individuals with LADA who tested positive for IA-2₍₂₅₆₋₇₆₀₎ and those who tested positive for other antibodies (FIG. 2).

Diagnostic challenges in LADA

Ideally, in order to identify LADA, positivity for islet autoantibodies should be tested in all patients diagnosed with T2DM. In fact, in clinics where GAD autoantibody testing is routinely carried out, the median time to insulin treatment for patients testing positive for GAD autoantibodies is markedly shorter than in clinics that do not offer GAD autoantibody testing¹³. As discussed in a later section, the choice of the most convenient therapy, especially in patients with a high GAD autoantibody titre, might have relevant advantages in terms of preserving β -cell function^{71,79}. Considering the prevalence of LADA and the increasing attention given to personalized therapy, testing for GAD autoantibodies could be considered for use as a routine test in patients newly diagnosed with T2DM. However, it is important to recognize that in relation to the achievable benefits, this procedure would be expensive if carried out consistently.

To date, islet autoantibodies are usually tested for only in patients thought to have T2DM if the clinical odds of the patient actually having LADA are high, which is generally assessed on the basis of BMI. Thus, adults with T2DM who have a normal weight are considered as potentially having LADA, whereas those who are overweight or obese are presumed to have T2DM and do not routinely undergo tests for autoantibodies. However, this approach does not consider the several studies showing that LADA can also be diagnosed in individuals with T2DM who are overweight or obese^{6,19,21,22}.

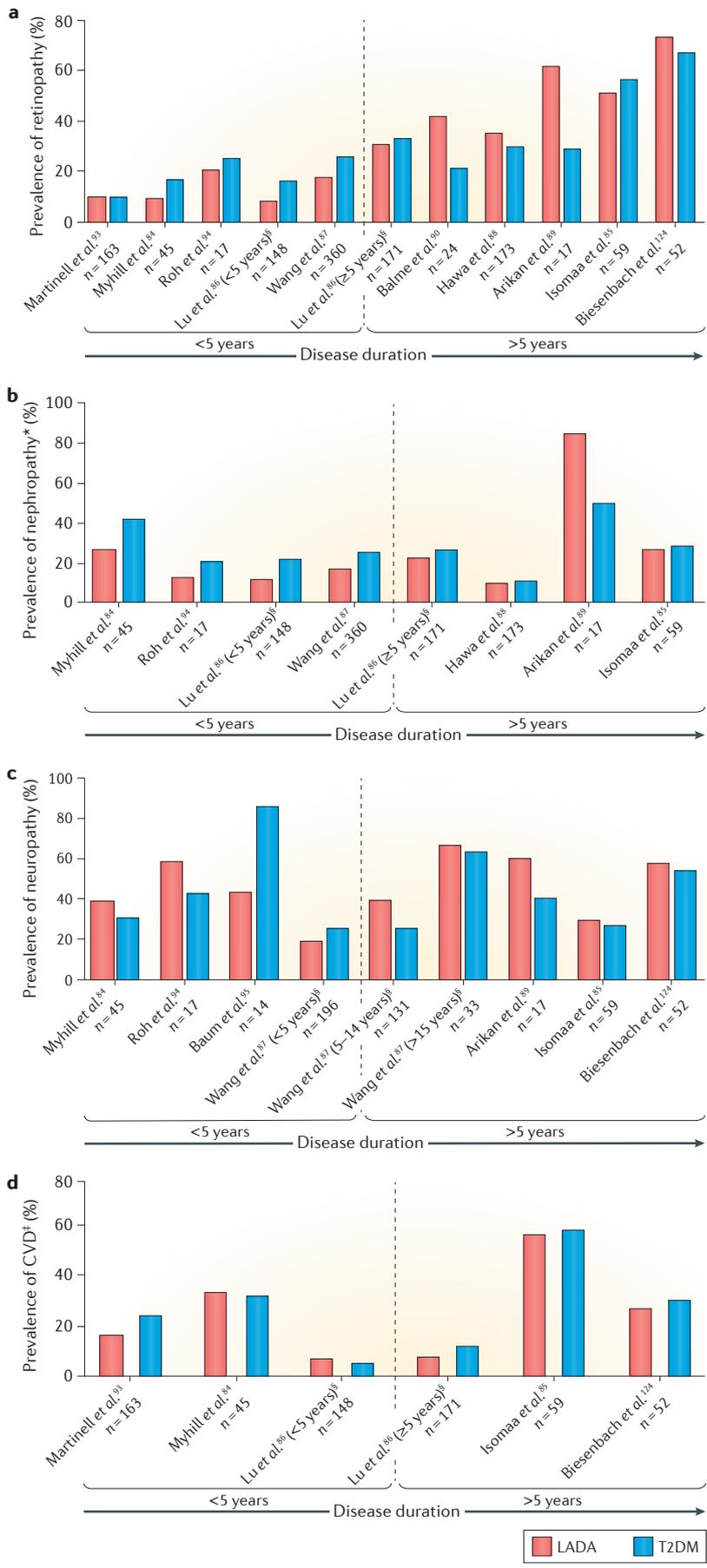
To identify patients with diabetes who have the highest odds of testing positive for autoantibodies, a screening tool⁸⁰ was developed to identify LADA in the clinical setting. In the retrospective phase of a two-stage study, five clinical parameters were found to occur more frequently in LADA than in T2DM: age of onset <50 years; acute symptoms before diagnosis (polydipsia, polyuria and unintentional weight loss); BMI <25 kg/m²; personal history of other autoimmune diseases; and a family history of autoimmune disease. In the second prospective phase, adults with newly diagnosed diabetes were enrolled to determine if a 'LADA clinical risk score' based on the five aforementioned parameters could identify LADA. The presence of at least two of these clinical features had 90% sensitivity and 71% specificity for identifying LADA, and the negative predictive value for a LADA clinical risk score ≤ 1 was 99%.

Risk of complications

The great burden of diabetes mellitus is mostly caused by the development of diabetes-related complications⁸¹. Diabetic ketoacidosis is the major acute complication and occurs in the context of absolute insulin deficiency. Therefore, this complication is generally rare in individuals affected by adult-onset autoimmune diabetes. Indeed, individuals with LADA, the most frequent form of adult-onset autoimmune diabetes, have detectable C-peptide levels at the time of diagnosis and show a slower decline in β -cell function than patients with T1DM⁴. Conversely, chronic complications (for example, cardiovascular disease, nephropathy, retinopathy and neuropathy) represent the major cause of decreased quality of life as well as increased morbidity and mortality in adults with diabetes mellitus⁸¹⁻⁸³. Although the different clinical and metabolic features of T1DM, LADA and T2DM suggest that they have different pathophysiologies and prevalences of chronic complications, few studies (most of which were underpowered) have investigated the risk of complications in LADA, and even fewer studies on this topic had a longitudinal study design^{84,85}.

Microvascular complications

Most studies that have compared the risk of microvascular complications between LADA, T2DM and T1DM were limited by a small sample size and by heterogeneity among the individuals enrolled with respect to ethnicity, metabolic control and disease duration, the last of which is the most relevant confounder. Indeed, an analysis of longitudinal data from the Fremantle Diabetes Study⁸⁴ showed that microalbuminuria was less frequent in patients with a recent LADA diagnosis (mean disease duration of 4 years) than in those with T2DM. Moreover, GAD autoantibody positivity was associated with a 62% reduced risk of developing microalbuminuria during a follow-up period of 3.7-4.4 years. A cross-sectional study confirmed that the rates of nephropathy and retinopathy were lower in Chinese patients with LADA than in those with T2DM⁸⁶. However, this observation was evident only for individuals who had a disease duration <5 years⁸⁷, and this lower risk was not seen in



patients who had a disease duration >5 years (FIG. 3a,b). Accordingly, studies that evaluated patients with a disease duration >5 years showed that patients with diabetes who were GAD autoantibody positive had equal or even higher rates of microvascular complications than those who were negative for GAD autoantibodies⁸⁸⁻⁹⁰ (FIG. 3a,b).

Prospective data from the BOTNIA Study⁸⁵ also showed that a disease duration >5.5 years was associated with an increased risk of retinopathy in patients with LADA (FIG. 3a). The low rates of microvascular complications observed in patients with LADA who have a short disease duration could in part be explained by the fact that T2DM is usually diagnosed later in the history of the disease than LADA^{91,92}. In fact, individuals with newly diagnosed T2DM are exposed to the detrimental effects of hyperglycaemia on the small retinal and renal blood vessels long before diagnosis, whereas this phenomenon has not been shown in LADA. However, a large Swedish registry study did not report differences between LADA and T2DM with respect to the prevalence of diabetic retinopathy at disease onset, in contrast to previous results⁹³ (FIG. 3a). In addition, no differences in the prevalence of microvascular complications were found among patients from South Korea with LADA, T2DM and T1DM⁹⁴. Most of the studies that have investigated diabetic neuropathy have shown that neuropathy was generally more prevalent in LADA than in T2DM^{84,85,87,89,94}, with the exception of a study by Baum *et al.*⁹⁵ (FIG. 3c).

Macrovascular complications

Individuals with LADA have a healthier lipid profile and lower blood pressure values, are leaner and have less central adiposity than individuals with T2DM^{6,7,19,22}. Collectively, these factors suggest a lower risk of coronary heart disease, stroke and peripheral artery disease

Figure 3 | Prevalence of chronic diabetes-related complications in latent autoimmune diabetes in adults and type 2 diabetes mellitus. The y axes depict the prevalence (%) of chronic diabetes-related complications, including retinopathy (part a), nephropathy (part b), neuropathy (part c) and cardiovascular disease (CVD; part d), in patients with latent autoimmune diabetes in adults (LADA; red bars) versus patients with type 2 diabetes mellitus (T2DM; blue bars). The x axes show the studies in which the data were reported. The studies reported on the x axes are stratified by a disease duration threshold of 5 years (depicted with a dashed line). The prevalences of retinopathy and nephropathy are generally lower in LADA (red bars) than in T2DM (blue bars) when the disease duration is <5 years and are equal when the two diseases or higher in LADA than in T2DM when the disease duration is >5 years. Neuropathy is more prevalent in LADA than in T2DM. The prevalence of CVD does not differ between LADA and T2DM. *The rates of microalbuminuria and/or macroalbuminuria are reported. †The rates of coronary artery disease are reported. ‡The data from these studies were split according to years of disease duration and are depicted separately in the graph. The term ‘n’ refers to the number of patients with LADA enrolled in each study.

in patients with LADA than in those who have T2DM. However, the evidence published thus far has failed to support this hypothesis (FIG. 3d). In the BOTNIA Study⁸⁵, 56% and 5% of the patients with LADA enrolled had a positive history of coronary artery disease and stroke, respectively, after 13 years of disease. These rates were similar to those found in people with T2DM, which were 58% and 7%, respectively⁸⁵. In the same study, the overall mortality during a follow-up of 5.7 years was 18% in LADA and 20% in T2DM (the difference was not statistically significant), and the cardiovascular mortality was 7.4% in LADA and 12.4% in T2DM (the difference was again not statistically significant). The HUNT2 study showed that patients with T2DM who were positive for GAD autoantibodies had a similar risk of all-cause mortality, cardiovascular disease and ischaemic heart disease as individuals with T2DM who were negative for GAD autoantibodies and an increased risk compared with healthy individuals⁹⁶. The Fremantle Diabetes Study⁸⁴ also did not show differences in the prevalences of cardiovascular disease and mortality between LADA and T2DM. Moreover, no differences in terms of cardiovascular outcomes were reported in a post-hoc analysis in the Collaborative Atorvastatin Diabetes Study⁸⁸.

Despite the healthier vascular risk profile of patients with LADA, the similar rates of cardiovascular disease in LADA combined with the evidence that the high mortality risk in autoimmune diabetes is not completely explained by hyperglycaemia⁸² suggest that a different pathophysiological mechanism underlies the development of cardiovascular disease in LADA compared with T2DM. This hypothesis warrants further investigation of the potential pathways mediating atherogenesis in autoimmune diabetes.

Treatment of LADA

Although different recommendations exist for the treatment of adults with T1DM and T2DM, no specific guidelines for the treatment of individuals with LADA have been published so far. As a consequence, the treatment of adults with LADA is currently guided by the clinical intuition and expertise of the physician. Owing to both frequent misdiagnosis and a lack of standard treatment options for LADA, most patients with LADA are initially treated with therapies intended for non-autoimmune forms of diabetes mellitus. This approach might result in rapid progression to an insulin-dependent state, especially in patients who are young and lean and who have high GAD autoantibody titres⁷⁷¹. Nevertheless, several studies have been carried out to test different treatment strategies (TABLE 3).

Addressing insulin deficiency

As a form of autoimmune diabetes, LADA is characterized by loss of β -cell mass and function, a crucial pathological event that should be managed by therapeutic strategies. Insulin treatment is the most straightforward therapy for replacing the low levels of endogenous insulin caused by β -cell loss. Interventions intended to preserve β -cell function have also been tested in LADA.

Insulin therapy. One of the main unanswered questions about the therapeutic strategies that should be undertaken in patients with LADA is if insulin therapy is really needed as an initial therapy for this form of autoimmune diabetes. Most of the studies investigating this issue were systematically reviewed by the Cochrane Collaboration in 2011, which showed that treatment with sulfonylureas was associated with worse metabolic control than insulin treatment, with earlier progression towards an insulin-dependent state^{79,97–102}. Therefore, it was recommended that sulfonylureas be avoided in favour of early insulin treatment. However, this recommendation relies on limited evidence⁹⁷ and does not take into account the need for a personalized medicine approach, which is recommended by the current international guidelines for the treatment of diabetes mellitus in adults¹⁰³. Our group has shown that progression towards an insulin-dependent state in LADA differs based on clinical and biochemical features⁷⁷¹. Patients with high GAD autoantibody titres have the highest risk of early insulin dependence. Moreover, a later onset of LADA is associated with a lower risk of needing insulin therapy. This observation raises the question of if early insulin therapy is the best approach in patients with LADA who have a low GAD autoantibody titre, particularly in older patients, who are at the highest risk of suffering from hypoglycaemia-related adverse effects.

Drugs to preserve β -cell function in LADA. The control of blood levels of glucose should not be the sole aim of LADA treatments; the preservation of β -cell function by suppressing the autoimmune destruction of β cells and by stimulating β -cell regeneration should also be pursued, taking advantage of the slow progression of LADA^{104,105}. Indeed, preserved insulin-secreting capacity is associated with better clinical outcomes and a decreased risk of complications^{106,107}.

Immune intervention trials have also been carried out for LADA and have shown some efficacy in preserving stimulated C-peptide levels and in achieving improved glycaemic control^{108,109}. Vaccination with GAD65 in LADA has shown a good safety profile during a 5-year follow-up period¹¹⁰. The tested molecules, however, are not currently available on the market.

Intriguingly, data suggest that drugs already approved for the treatment of T2DM might be useful for the treatment of LADA. Preclinical data from animal models have shown that treatment with glucagon-like peptide 1 (GLP1) induces β -cell self-renewal, reduces β -cell apoptosis and promotes β -cell neogenesis from ductal cell precursors¹¹¹. However, *in vitro* data from human cells have supported only a possible protective effect of incretin hormones on human β cells, specifically in terms of reducing apoptosis¹¹², and no randomized controlled trials testing the implementation of GLP1 receptor agonists in patients with LADA are available. A large prospective study that evaluated predictors of the glycaemic response to GLP1 receptor agonists found that patients diagnosed with T2DM who tested positive for GAD and/or IA-2 autoantibodies and who had low C-peptide levels

Table 3 | Studies investigating pharmacological therapies for LADA

Study design	Drugs tested	Study population	Main findings	Refs
Pilot randomized, open-label study	Insulin versus SU	<ul style="list-style-type: none"> • 10 participants with NIDDM • Positive for ICAs 	<ul style="list-style-type: none"> • Serum C-peptide response improved significantly within 6 and 12 months in the insulin group and decreased in the SU group. • Glycaemic control worsened in the SU group, but not in the insulin group. 	98
Randomized, multicentre, open-label study	Insulin versus SU	60 participants with diabetes, positive for GAD autoantibodies, disease duration ≤5 years	After a mean follow-up of 57 months, a lower rate of progression to an insulin-dependent state was observed in the insulin group.	99
Randomized, open-label study	Insulin versus diet with or without OHA (metformin and/or SU)	37 participants with NIDDM, positive for ICAs or GAD autoantibodies	<ul style="list-style-type: none"> • HbA_{1c} levels were stable in the insulin-treated group but increased in the diet ± OHA group. • No significant differences in C-peptide levels were observed between the two groups. 	100
Randomized, open-label study	Rosiglitazone with insulin versus insulin alone	23 participants with LADA, fasting C-peptide levels >0.3 nmol/l	After 12 and 18 months, measures of β-cell function were stable in the group receiving rosiglitazone with insulin but decreased in the insulin-alone group.	121
Phase II, randomized, placebo-controlled, dose-escalation clinical trial	Alum-formulated recombinant GAD65 (GAD65 vaccination)	47 participants with T2DM, GAD autoantibody positive	Fasting C-peptide levels remained stable for 5 years after drug administration in the groups treated with 4, 20 and 100 micrograms of the GAD65 vaccine but decreased in the placebo and 500-microgram groups.	108,110
Longitudinal observational study	GLP1-RA	620 participants with T2DM commencing a GLP1-RA	<ul style="list-style-type: none"> • GAD or IA-2 islet autoantibody-positive participants had a nonsignificant reduction in the adjusted mean level of HbA_{1c} (-4.6 mmol/mol (95% CI -10.3 to 1.1)), which was lower than the reduction observed in antibody-negative participants. • Antibody-positive participants experienced a 17% reduction in insulin dose (versus 40% in antibody-negative participants, <i>P</i> < 0.01). 	113
Randomized, open-label study	Sitagliptin and insulin versus insulin alone	30 participants with LADA	Measures of β-cell function evaluated after 12 months of treatment were stable in the group receiving sitagliptin and insulin but were decreased in the insulin-alone group.	115
Prespecified exploratory analysis of a randomized controlled trial	Linagliptin versus glimepiride	118 participants with LADA	Fasting C-peptide levels increased from baseline at weeks 28, 52 and 104 in participants treated with linagliptin but decreased in glimepiride-treated patients. No differences in metabolic control were found.	114
Post-hoc analysis of data pooled from five randomized, placebo-controlled, 24-week phase III studies	Saxagliptin	133 participants, GAD autoantibody-positive (98 in the saxagliptin arm and 35 in the placebo arm)	<ul style="list-style-type: none"> • Saxagliptin was effective in lowering blood glucose levels and was generally well tolerated in GAD autoantibody-positive participants. • Saxagliptin increased β-cell function in GAD autoantibody-positive participants. 	116

GAD, glutamic acid decarboxylase; ICA, islet-cell autoantibody; GLP1-RA, glucagon-like peptide 1 receptor agonist; IA-2, tyrosine phosphatase IA-2; LADA, latent autoimmune diabetes in adults; NIDDM, non-insulin-dependent diabetes mellitus; OHA, oral hypoglycaemic agent; SU, sulfonylurea; T2DM, type 2 diabetes mellitus.

exhibited a lower reduction in HbA_{1c} than patients who tested negative for autoantibodies and had high C-peptide levels¹¹³. This study suggests that autoantibody positivity and C-peptide levels could be useful for the stratification of patients with diabetes in terms of therapy response; individuals with LADA are less likely to benefit from incretins than patients with T2DM in terms of reductions in levels of HbA_{1c}. However, this study does not exclude the potential benefits of GLP1 receptor agonists in terms of β-cell protection, which is suggested by a mean 17% reduction in daily insulin dose, even in patients who are autoantibody positive and who have C-peptide levels <0.25 pmol/l. Furthermore, other studies have reported some benefits in terms of the preservation of C-peptide levels by using the dipeptidyl peptidase 4 inhibitors sitagliptin, linagliptin and saxagliptin^{114–116}.

Lifestyle factors. Smoking has been hypothesized to be associated with decreased autoimmunity and a larger pancreatic reserve, as suggested by a lower GAD autoantibody titre, higher C-peptide levels and a reduced risk of developing LADA in smokers than in non-smokers^{117,118}. However, data from larger populations have called this hypothesis into question¹¹⁹. Moreover, an increase in daily alcohol intake has been associated with increased GAD autoantibody levels; however, further longitudinal studies are needed to better investigate the possible association between alcohol consumption and autoimmunity in adults with diabetes¹²⁰.

Addressing insulin resistance

In addition to insulin deficiency, insulin resistance is also a clinical feature of LADA, although it is less severe than in T2DM. Thus, patients with LADA who have

a more pronounced insulin-resistant phenotype might benefit from the addition of insulin sensitizers to insulin therapy. However, no randomized controlled trials have tested metformin in LADA; only rosiglitazone has been tested as an insulin sensitizer in LADA, with encouraging results¹²¹, demonstrating the potential use of peroxisome proliferator-activated receptor γ agonists.

Clinical outlook

In summary, as LADA has clinical features that are midway between those of T1DM and T2DM, treatments aimed at both preserving and restoring β -cell function by suppressing autoimmunity and stimulating β -cell proliferation as well as at improving insulin sensitivity and ameliorating glycaemic control are warranted. Accordingly, the use of combination therapies

to target different pathways could be the best choice, but large randomized controlled trials will be required to confirm this hypothesis¹⁰⁶. Based on the current evidence, an early start to insulin treatment together with lifestyle interventions (diet and physical activity) could be advised for patients with LADA, especially in younger patients with high GAD autoantibody titres, because such treatment modalities are able to both control hyperglycaemia and preserve β -cell function by reducing β -cell stress and preventing glucotoxicity^{121,122}. Other agents, such as gliptins or insulin sensitizers, could be combined with insulin based on patients' clinical features, such as BMI, HbA_{1c} levels and C-peptide levels. Finally, it should always be remembered that patients with diabetes mellitus, independent of subtype, need a multifactorial approach to treatment that takes into account the complexity of the patient in terms of comorbidities¹²³.

Conclusions

Adult-onset autoimmune diabetes encompasses a wide spectrum of heterogeneous clinical and metabolic phenotypes, ranging from classic T1DM with onset in adult life to LADA^{6,19,21,22,25}. If we accept the definition whereby LADA can be identified by positivity for any islet autoantibody (not necessarily GAD autoantibodies)¹², we have to recognize that this form of diabetes mellitus also has a considerable degree of heterogeneity. This heterogeneity encompasses not only patients who have clear signs of insulin deficiency, which is usually associated with strong markers of autoimmunity²⁹, but also patients who have a clinical phenotype that resembles T2DM but who are still positive for fairly weak markers of autoimmune reactivity⁵⁷. Therefore, the heterogeneity of LADA could represent a progressive phenotypic spectrum between the two most common forms of diabetes mellitus, T1DM and T2DM⁷⁵ (FIG. 4). This hypothesis challenges the conventional perception that immune-mediated processes are not relevant to the pathogenesis of T2DM. Further studies are warranted to clarify if the low-grade inflammation that characterizes visceral obesity is linked to an autoimmune process that could influence the understanding of the pathogenesis of T2DM⁵⁷.

The difficulties in comparing the results of different studies investigating the microvascular and macrovascular complications associated with adult-onset autoimmune diabetes are partly attributable to the different criteria used to screen patients and to the heterogeneity of the clinical phenotypes of the enrolled patients^{81,84–94}. Overall, studies with larger cohorts and a longitudinal design are needed to investigate both the pathophysiology and the epidemiology of the complications associated with adult-onset autoimmune diabetes.

We have not yet identified the best therapy for LADA. The identification of efficacious therapies for patients with LADA has been limited by the great degree of variability of the biochemical and clinical features at clinical presentation during patient screening for intervention trials, which makes it difficult to assess the response to therapy^{79,97–123}. Defined categories of disease are

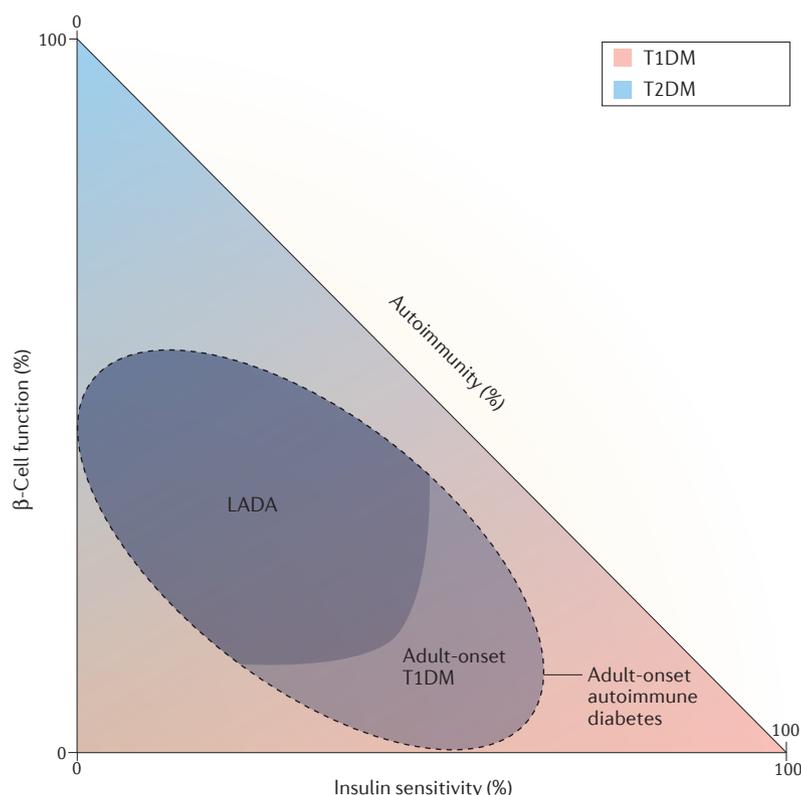


Figure 4 | Interactions between β -cell function, autoimmunity and insulin sensitivity.

β -Cell function, autoimmunity and insulin sensitivity are three key pathological factors that are involved in the pathogenesis of diabetes mellitus. The spectrum of adult-onset diabetes mellitus encompasses all three forms of diabetes, namely, type 1 diabetes mellitus (T1DM), type 2 diabetes mellitus (T2DM) and adult-onset autoimmune diabetes, without defined limits. The figure depicts the continuum (0–100%) of continuously distributed pathological factors and defines the categories of diabetes mellitus according to these variables. Classic T2DM (upper left corner) is characterized by the absence of autoimmunity, well-preserved β -cell function and low insulin sensitivity. Classic T1DM (lower right corner) is characterized by severe autoimmunity, loss of β -cell function and preserved insulin sensitivity. Along the continuum, adult-onset autoimmune diabetes appears midway between the features of T1DM and T2DM, making it difficult to draw defined limits between the different forms of diabetes (dashed lines). Latent autoimmune diabetes in adults (LADA) accounts for the majority of the cases of adult-onset autoimmune diabetes and shows pathological features closer to those of T2DM than to those of adult T1DM, which is more similar to classic T1DM.

commonly needed to offer guidance for the choice of treatment. However, it is difficult to define disease categories for adult-onset autoimmune diabetes owing to the continuous distribution of clinical and metabolic variables (such as autoimmunity, β -cell function and insulin sensitivity). The existence of diverse metabolic

and clinical phenotypes in adult-onset autoimmune diabetes makes it difficult to establish an *a priori* treatment algorithm and therefore draws attention to the concept of a personalized medicine approach to therapy that takes into account the intrinsic characteristics of each patient.

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