Obesity, metabolic syndrome, and type-2 diabetes mellitus are 3 interrelated conditions that share a number of pathophysiological mechanisms and that are frequently observed to lead, in succession, to cardiovascular complications. The fact that their prevalence is increasing alarmingly should prompt all healthcare professionals urgently to implement measures to prevent these complications. The most effective, though also the least adopted, are those related to lifestyle modification. Drug treatment targeted at controlling risk factors (eg, hypertension, dyslipidemia, and thrombophilia), metabolic abnormalities, and excess weight is also necessary.

Key words: Diabetes. Metabolic syndrome. Obesity.

INTRODUCTION

Few will disagree that obesity is fast becoming the 21st century’s most serious health problem. It is a condition which begins in childhood, flourishes in adulthood, and leads to multiple health problems. Genetic and molecular factors, other exacerbating circumstances, and environmental and behavioral triggers all play a part in its development and determine its treatment. Type 2 diabetes mellitus (DM2) is a metabolic disease arising from flaws in the action of insulin in the peripheral tissues. The resulting chronic hyperglycemia is accompanied by damage to organs, particularly the eyes, kidneys, and heart, as well as to nerves and arteries.

Obesidad, síndrome metabólico y diabetes: implicaciones cardiovasculares y actuación terapéutica

Obesidad, síndrome metabólico y diabetes mellitus tipo 2 son tres enfermedades interrelacionadas que compar- ten mecanismos de aparición y evolución y con frecuen- cia se van combinando sucesivamente y ocasionan com- plicaciones cardiovasculares. Su prevalencia crece alarmantemente y debería impeler a los profesionales de la salud y a los gestores a implantar medidas urgentes para prevenir la aparición de complicaciones. Las más efícares, aunque menos practicadas, son las relaciona- das con el estilo de vida. También son necesarios trata- mientos farmacológicos destinados al control de los fac- tores de riesgo (hipertensión, dislipemias, trombofilía), las alteraciones metabólicas y el propio exceso de peso.

Palabras clave: Diabetes. Síndrome metabólico. Obesi- dad.

DEFINITIONS

Obesity

Obesity is defined as an excess of corporal adiposity. For practical reasons, body weight has been used as an indirect measure of the degree of adiposity as the latter is not easy to measure with the usual tests. In the eighties, the concept of body mass index (BMI) was introduced and cut points were established to define overweight and obesity in adult men and women. Values were later established for children and adolescents (Table 1).

Independent of the fact that high BMI values are associated with adverse risk profiles for morbidity and mortality, particularly in relation to DM2 and
Atherothrombotic cardiovascular disease (CVD), various sub-types of obesity have been described which complement the apparent dose-response relationship between BMI and its impact on health. The most classic sub-types are android obesity (abdominal) and gynoid obesity (gluteal obesity). Less well-known sub-types are, however, also of interest. For example, a phenotype corresponding to metabolically obese normal-weight (MONW) individuals has been reported. These individuals have a normal BMI together with some of the characteristics of obese individuals such as insulin resistance, central adiposity, low levels of high density lipoproteins (HDL) and high levels of triglycerides, and high blood pressure (HBP). At the same time, metabolically healthy obese (MHO) individuals have also been identified. These individuals have BMI>30, but none of the metabolic alterations which are typical of obese individuals.

The existence of these 2 “paradoxical” sub-types led to several studies over the last 2 decades which aimed to metabolically characterize and quantify visceral adipose tissue as opposed to subcutaneous adipose tissue. These studies showed that the quantity of visceral adipose tissue is directly correlated in both men and women with a seriously altered metabolic risk profile which precedes the development of DM2 and cardiovascular disease. Therefore, although it is true that obesity increases the risk of chronic disease, it seems clear that patients with visceral obesity form a sub-group of individuals with the most serious metabolic alterations. It has been demonstrated, for instance, that the regional accumulation of fatty tissue in the visceral deposits is a more reliable predictor of cardiovascular risk than the total quantity of body fat. The multi-country INTERHEART case-control study confirmed the importance of adiposity, and particularly abdominal adiposity, as a risk factor for acute myocardial infarction.

Metabolic Syndrome

In recent years, MS has caused an authentic revolution in vascular risk stratification and has become one of the most widely used concepts within the health sciences. It describes the accumulation within an individual patient of several very prevalent conditions in western countries, all of which are risk factors for CVD and DM2. The 2 underlying mechanisms of MS are abdominal adiposity and insulin resistance, it is exacerbated by age, genetic conditioning, and an inappropriate lifestyle which includes physical inactivity and the consumption of high calorie, high-fat foods, concentrated carbohydrates, and salt. These metabolic alterations occur simultaneously more frequently than would be expected by chance and the concurrence of several factors increases cardiovascular risk over and above the risk associated with the individual factors alone.

There are numerous definitions of MS. Although all of the classifications include the syndrome’s essential components, they differ on details or on whether to include components which are difficult to measure. The two most widely used definitions are those of the International Diabetes Federation classification and the definition included in the latest revision of the ATP-III classiﬁcation and the definition included in the latest revision of the ATP-III (Adult Treatment Panel III of the United States’ National Cholesterol Education Program). The 2 classifications are very similar and are summarized in Table 2.

The risk factors constituting MS are atherogenic dyslipidemia, high blood pressure, glucose metabolism disorders, and proinflammatory and prothrombotic states, ie all of the classic CVD risk factors except for smoking, family history, and age. Atherogenic dyslipidemia includes higher levels of apolipoprotein B containing lipoproteins and triglycerides, a greater number of small low density lipoprotein (LDL) particles (with a higher atherogenic cholesterol content), and low concentrations of HDL. Although they are not included in the definition of MS, other biological markers can sometimes be useful, such as those related to adipose tissue (percentage of abdominal fat using digital tomography, leptin/adiponectin ratio), other markers of dyslipidemia (apolipoprotein B or size of LDL particles), endothelial dysfunction (measured by the vasodilatory response in the humeral artery), inflammation markers (C-reactive protein) or thrombosis markers (high fibrinogen or plasminogen activator inhibitor–1) (Table 3).

### Abbreviations

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Definition</th>
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<tbody>
<tr>
<td>BMI</td>
<td>body mass index</td>
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<tr>
<td>CVD</td>
<td>cardiovascular disease</td>
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<tr>
<td>DM2</td>
<td>type 2 diabetes mellitus</td>
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<tr>
<td>HBP</td>
<td>high blood pressure</td>
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<tr>
<td>HDL</td>
<td>high density lipoproteins</td>
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<tr>
<td>MS</td>
<td>metabolic syndrome</td>
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</table>

### Table 1. Definitions of Obesity According to Body Mass Index

<table>
<thead>
<tr>
<th>Category</th>
<th>Children and Adolescents</th>
<th>Adults</th>
</tr>
</thead>
<tbody>
<tr>
<td>Underweight</td>
<td>&lt;p5</td>
<td>&lt;18.5</td>
</tr>
<tr>
<td>Normal weight</td>
<td>p5-85</td>
<td>18.5-24.9</td>
</tr>
<tr>
<td>Risk of overweight</td>
<td>p85-90</td>
<td></td>
</tr>
<tr>
<td>Overweight</td>
<td>&gt;p95</td>
<td>25-29.9</td>
</tr>
<tr>
<td>Grade 1 obesity</td>
<td>—</td>
<td>30-34.9</td>
</tr>
<tr>
<td>Grade 1 obesity</td>
<td>—</td>
<td>35-39.9</td>
</tr>
<tr>
<td>Morbid obesity</td>
<td>—</td>
<td>40-49.9</td>
</tr>
<tr>
<td>Super-morbid obesity</td>
<td>—</td>
<td>&gt;50</td>
</tr>
</tbody>
</table>

*pBMI percentile by age group (Centers for Disease Control and Prevention), a Body mass index (BMI) = weight (in kg)/height (in m) squared (NIH and OMS), p = percentile.*
**TABLE 2. Diagnostic Criteria for Metabolic Syndrome**

<table>
<thead>
<tr>
<th>Condition</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Central adiposity</td>
<td>Waist circumference for European population: males, &gt;94; females, &gt;80 cm and 2 or more of: Triglycerides &gt;150 mg/dL or specific treatment; HDL&lt;40 (males) or &lt;50 mg/dL (females) or treatment; Blood pressure &gt;130/85 mm Hg or treatment for hypertension; Baseline blood glucose &gt;100 mg/dL or prior diagnosis of diabetes</td>
</tr>
<tr>
<td>National Cholesterol Education Program (ATP-III) (2002)</td>
<td>Three or more of the following criteria: Fasting glucose &gt;110 mg/dL [100]; Blood pressure &gt;130/85 mm Hg [or treatment]; Triglycerides &gt;150 mg/dL [or treatment]; HDL cholesterol &lt;40 mg/dL in males; &lt;50 mg/dL in females; Abdominal adiposity Waist &gt;102 cm (males); Waist &gt;88 cm (females)</td>
</tr>
</tbody>
</table>

HDL indicates high density lipoproteins.
The details in square brackets were added in 2005 to adapt to the IDF definition.

**Diabetes**

The American Diabetes Association diagnostic criteria for diabetes are clear\(^\text{14}\): \(a\) casual plasma glucose level >200 mg/dL in a patient with characteristic symptoms ("casual" is defined as an analysis performed at any time of day independently of the time of the last food intake), polyuria, polydipsia, and weight loss; or \(b\) fasting glucose level >126 mg/dL confirmed in 2 determinations (fasting is defined as the absence of caloric intake during at least 8 h); or \(c\) plasma glucose values 2 h after an oral glucose load >200 mg/dL; the test should be performed according to the WHO guidelines, using a standard drink containing the equivalent of 75 g of anhydrous glucose dissolved in water.

As well as diabetes, other pre-diabetic syndromes which are reflected in the concept of glucose metabolism disorders are diagnosed in patients who do not meet the criteria for a diagnosis of diabetes but who have higher than normal basal glucose levels or after loading. Figure 1 summarizes the criteria for the diagnosis of altered basal glucose values and glucose intolerance.\(^\text{14}\)

**PHYSIOPATHOLOGICAL ASPECTS**

Details on the mechanisms underlying the type of vascular injuries which are produced by the 3 inter-related conditions under consideration, ie obesity, MS, and DM2 go beyond the scope of this review article. Few brief pointers are provided.

Obesity is a risk factor for atherosclerosis due to a variety of mechanisms which are summarized in Figure 2. Insulin resistance, which is the common link with MS, leads to glucose metabolism disorders, dyslipidemia, high blood pressure, endothelial dysfunction, and inflammation induced by cytokine imbalance. The latter, ie an excess of harmful cytokines (interleukins 6 and 18, tumor necrosis factor alpha, and leptin being among the most important) combined with a scarcity of protector cytokines (adiponectin).\(^\text{15}\)

Central adiposity, one of the principal characteristics of MS, provides the foundation for the portal/visceral theory, which suggests that the increase in adiposity, particularly in visceral deposits, leads to an increase in the flow of free fatty acids and the inhibition of insulin action. The large quantity of non-sterified fatty acids reduces the use of glucose by skeletal muscle, stimulates the hepatic production of very low-density lipoproteins and glucose, and propitiates the acute secretion of insulin.

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**Figure 1.** Concept of glucose metabolism disorder (American Diabetes Association criteria\(^\text{11}\)). Diabetes is diagnosed when several determinations of basal glucose levels (after 12 h of fasting) exceed 126 mg/dL or if glucose levels 2 h after an oral load of 75 g of glucose (OGL) exceed 200 mg/dL. Glucose levels of 100 to 126 mg/dL indicate a diagnosis of “abnormal glucose levels” (AGL) and between 140 and 200 mg/dL after OGL indicate “glucose intolerance” (GI).
The lipotoxic effect in the pancreatic beta cells caused by free fatty acids over the long term could be part of the connection between obesity, insulin resistance, and the development of DM2. However, the portal theory has been placed in quarantine and there is some evidence that it is unable to explain, by itself, all of the metabolic anomalies present in abdominally obese individuals.\footnote{16,17}

There was considerable debate as to whether MS increases the risk of cardiovascular disease to a greater degree than the sum of its individual parts, though it has now clearly been shown that this is the case.\footnote{16} In the first place, epidemiological studies have demonstrated that the degree of risk involved when multiple risk factors are present is greater than the sum of the risks attributable to each individual factor; in other words, under these circumstances, risk increases geometrically rather than linearly. In the second place, the algorithms habitually used to calculate cardiovascular risk do not include several metabolic factors which have been shown to independently increase the risk of vascular complications. These include the prothrombotic state, inflammatory status, and hypertriglyceridemia. The additional risk associated with these factors exceeds that attributed to conventional risk factors. In the third place, some of the risk attributed to conventional factors probably derives from factors which have not been considered or measured. Finally, as MS progresses and often culminates in DM2, the risks associated with the latter should be added to those already mentioned. The evolution of MS in terms of its pathological consequences are shown in Figure 3.\footnote{16}

Figure 4 summarizes the latest theories regarding the relationship between obesity and MS, which is posited as a dysfunction of adipose tissue with insulin resistance appearing later when compensatory response mechanisms have been exhausted.\footnote{17}

From what has been described here, it is clear that MS is a first order risk factor for atherothrombotic complications. Its presence or absence should therefore be considered an indicator of long-term risk. On the other hand, over the short term (5-10 years) risk is better calculated using the classical algorithms (Framingham, SCORE, REGICOR), as they include age and sex, total cholesterol or LDL, and smoking.\footnote{16}

The basic cardiovascular complications produced by hyperglycemia in DM2 are microangiopathic disease (diabetic nephropathy, retinopathy, and diabetic neuropathy) and CVD (diabetic macroangiopathy). These include coronary heart disease and stroke as the main complications, although peripheral arterial disease is also common and tends to be more serious (higher indices of serious ischemia and amputations) when it is cause by diabetes than when it is caused by other mechanisms. Cardiovascular diseases is the most important complications of DM2, both in clinical and numerical terms and represent up to 80\% of the macrovascular complication attributed to the disease.\footnote{18,19} Apart from hyperglycemia, other risk factors (notably high blood pressure, dyslipidemia, and obesity) are also very prevalent in diabetic patients.

Micro- and macroangiopathy exist side by side and their harmful effects are multiplied in DM2. Damage to the endothelium produced by accumulation of glucosilation sub-products, reduced production of nitric oxide, and endothelial inflammation are among the most characteristic mechanisms involved in the process.

**SOME EPIDEMIOLOGICAL DATA**

Over the last 50 years, people have experienced drastic changes in their environment, behavior, and lifestyles.
These changes have led to the alarming increase in incidence of diabetes and obesity. The substitution of manual labor by mechanized processes, increased consumption of “empty calories,” the globalization of technology, and improvements in transport are some of the factors which have led to this pandemic. One of the most recent studies into the prevalence of overweight, which was carried out by the WHO and the International Obesity Task Force, found that the worldwide prevalence, after adjusting for ethnic differences, was 1700 million overweight individuals, of which 312 million were obese. The trend in the obesity figures means that it is now considered one of the greatest health threats in the developed world. Data from successive cohorts of the National Health and Nutrition Examination Survey (NHANES) show that the mean waist circumference of adults in the USA increased by 3.1 cm between the 1988-1994 and 1999-2000 surveys. After adjusting by age, the largest increases in waist circumference were observed in the youngest age group (20-29 years). Over the past 20 years, prevalence rates for obesity have tripled in developing countries which have adopted western lifestyles. These have shown the typical patterns of reduced physical activity and over consumption of high energy-density meals. The last survey into the prevalence of overweight in CVD by the European Society of Cardiology (Euro-Heart Survey) was carried out in 8547 patients with chronic ischemic heart disease and found a prevalence of overweight (BMI>25) of 83% and of obesity (BMI>30) of 38%. Overall, 55% of the sample had central obesity (waist circumference >102 cm in men, >88 cm in women) (unpublished data).
There have been several studies into the prevalence of obesity in the Spanish general population. For example, Figure 5 shows data from the MESY AS register obtained from 18,778 healthy members of the working population (78% male).23

One of the reasons for the current interest in MS is its high prevalence. In the Spanish MESYAS register of healthy members of the working population, 12% were found to have the condition (Figure 6).23 This compares to figures of 22% for the Spanish general population, 41% for patients with heart disease in Spain, and 24% of those included in the United States NHANES survey.22

Finally, the DM2 epidemic is expanding principally in developed and developing countries. Estimates are not very encouraging and have taken public health authorities by surprise. Worldwide, the number of those with DM2 is expected to grow from the current 150 million to 220 million in 2010 and to 300 million in 2025.24 The great majority will have DM2 associated with obesity and a sedentary lifestyle. This trend towards an increased prevalence of DM2 and obesity presents a real challenge for public health systems practically the whole world over and will continue to do so into the foreseeable future (Figure 7).

Figure 5. Prevalence of weight alterations in men (A) and women (B) in the MESYAS register of 18,778 healthy men and women.

Figure 6. Prevalence of metabolic syndrome in the MESYAS register by age and gender in 18,778 healthy members of the working population.
DIAGNOSTIC APPROACH

Obesity and Metabolic Syndrome

Diagnosis in individuals with obesity and MS should take into account the progressive nature of these conditions. In general, their evolution can be summarized in 4 phases, beginning with the presence of certain causal factors (central adiposity and insulin resistance, as well as genetic factors). These give rise to the metabolic alterations described above (high blood pressure, dyslipidemia, and glucose metabolism disorders) which in turn are followed by the development of vasculopathy. This is initially subclinical but eventually manifests as characteristic atherothrombotic complications and DM2 (Figure 3). Diagnosis should therefore not be limited to determining the presence of metabolic alterations, but should also explore the possibility of vasculopathic complications.

Given the high risk associated with both diseases over the long term, it would seem logical to suggest that overall cardiovascular risk be assessed in all obese individuals or those with MS. Such an assessment should include checking on the presence and magnitude of MS components. Medium-term risk should be evaluated using risk prediction models adapted whenever possible to the local population, and the possibility of sub-clinical vasculopathy should be investigated using the appropriate procedures.

Signs of asymptomatic vasculopathy can be studied in all in vascular territories mostly affected by atherosclerosis (Table 4). If coronary heart disease is suspected, stress tests should be used to detect myocardial ischemia or alterations in perfusion together with non-invasive coronary angiographic imaging via multislice computed tomography or the classical technique. The possibility of an effect on cerebral blood flow can be studied by determining the intima media thickness of the common carotid artery, while the impact on peripheral circulation is measured most precisely using abdominal ultrasound to determine the diameter of the aorta. The ankle-brachial index can also be used as a non-invasive method to detect peripheral vascular disease, which correlates with vascular risk. Other, more general signs of sub-clinical vasculopathy are microalbuminuria and erectile dysfunction.

Diabetes

Diabetes is now considered a vascular disease even when it clinical complications are not present. This approach is clearly important from the treatment point of view, though it is more doubtful when it comes to diagnosis, and has led to disagreement with the European Society of Cardiology’s clinical practice guidelines on diabetes and pre-diabetes which strongly recommend that tests to detect myocardial ischemia are performed on all patients with DM2.

<table>
<thead>
<tr>
<th>Territory</th>
<th>Test</th>
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</table>
| Coronary  | Tests of ischemia  
|           | Multislice computed tomography  
|           | Coronary angiography |
| Encephalic| IMT carotid   |
| Periphery | Abdominal ultrasound |
| General   | Ankle/brachial index |
|           | Albuminuria |
|           | Erectile dysfunction |
|           | Endothelial function |

IMT indicates intima media thickness.

Figure 7. Diabetes cases (in millions of individuals) in 2000 and predicted for 2030. 

TABLE 4. Study of Sub-Clinical Vasculopathy
Table 5 provides a summary of the general approach to treating the metabolic syndromes. The primary objective is to prevent onset of CVD, though it is equally important to delay or avoid onset of DM2 if it is not already present. Life-style

Lifestyle is key to the treatment of DM2, MS, and obesity. If all of these conditions are due to the expression of a particular genetic predisposition exacerbated by an inappropriate lifestyle, then the first step in treatment is to modify that lifestyle. The three basic components in any such improvement are diet, exercise, and behavior.

The first essential step is weight loss (Figure 5). This apparently simple step (which is in fact the most difficult to achieve and, therefore, the least frequently achieved) has demonstrated numerous beneficial effects: it reduces mortality, improves lipid profiles, reduces blood pressure and left ventricle mass in hypertension, and it reduces inflammation markers in DM2. Furthermore, in patients with MS it has been shown that it slows progression to DM2. The same beneficial effects have also been observed in obese patients.

As well as the quantitative aspect of a reduction in caloric uptake, dietary composition can also affect vascular health. Avoiding fats other than those from olive oil, increasing the consumption of fruits, vegetables, and whole-food cereals, avoiding industrially sweetened products and fatty meats, and eating white and especially blue fish are some of the recommendations in this regard (Table 6).

Exercise has similar beneficial effects in MS, obesity, and DM2. As well as helping to achieve and maintain weight loss, it also reduces all of the components of MS and reduces the incidence of CVD.

It may be stating the obvious to insist that patients should be encouraged to give up smoking, but it is not enough only to offer advice; health care professionals should ensure they provide the requisite support for stopping smoking at all levels of cardiovascular risk.

Treatment of High Blood Pressure

According to the ADA and ESH/ESC guidelines, blood pressure targets in diabetes mellitus should be <130/80 mm Hg. The same values are recommended for patients with MS.

Angiotensin converting enzyme (ACE) inhibitors have demonstrated multiple beneficial effects on cardiovascular complications and for that reason are considered the first line, or preferred treatment for high blood pressure, particularly when established or incipient kidney disease is present. Some well-known studies have also shown that calcium antagonists are effective against CVD. In DM2 patients, either of these treatments could be considered the most appropriate choice, alone or in combination. Beta-blockers are indicated where coronary artery disease or heart failure are present. Diuretics showed a satisfactory anti-hypertensive effect in some studies, though they also led to a worsened metabolic status and have no effect on arterial rigidity. For these reasons, beta-blockers and diuretics are considered complementary to ACE inhibitors and calcium channel blockers, with which they can be carefully used in combination to achieve appropriate levels of blood pressure control.

Lipid-Lowering Treatment

Lipid-lowering treatments are used based on the initial levels of lipids in blood with the aim of reaching targets set in international guidelines (Figure 8), ie LDL<100 mg/L, though a value of <70 mg/dL may be selected if

<table>
<thead>
<tr>
<th>TABLE 5. Overview of Treatment for Metabolic Syndrome, Obesity, and Diabetes</th>
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<tr>
<td><strong>Aspect</strong></td>
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<tr>
<td>Essential</td>
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<tr>
<td>Complementary</td>
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<tr>
<td></td>
</tr>
<tr>
<td>Adjunctive drugs</td>
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<tr>
<td>Complications</td>
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<td></td>
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<tr>
<td></td>
</tr>
<tr>
<td>Revascularization</td>
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</table>

<table>
<thead>
<tr>
<th>TABLE 6. Dietary Components and Vascular Risk</th>
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<tbody>
<tr>
<td><strong>Harmful</strong></td>
</tr>
<tr>
<td>Excess calories</td>
</tr>
<tr>
<td>Saturated fats</td>
</tr>
<tr>
<td>Salt</td>
</tr>
<tr>
<td>Alcohol (in excess)</td>
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</tbody>
</table>
cardiovascular risk is particularly high. A secondary, though very frequent and decisive objective in MS, obesity, and DM, is to lower cholesterol, other than HDL cholesterol, to levels <130 mg/dL (optional <100 mg/dL). In order to prevent pancreatitis, triglycerides become a primary target for treatment when they exceed 500 mg/dL.  

13,40 Statins are the drug of choice to lower LDL levels. They have been shown to reduce serious cardiovascular complications by 21% and 23% in diabetic and non-diabetic patients, respectively, in both primary and secondary prevention. 41 Statins are therefore essential treatment in patients with diabetes mellitus. The dose should be based on the size of the reduction required in LDL cholesterol. Any of the drugs in the group should be suitable; the important thing is to pursue and achieve the required dose and the targets set.

Fibrates are considered more effective than statins in lowering triglyceride and raising HDL levels, due to their agonist effect on PPARα. They might therefore appear to be an ideal treatment for atherogenic dyslipidemia in diabetes mellitus. 42 However, the FIELD study showed that fenofibrate, one of the most widely used fibrates, did not significantly reduce the risk of coronary events. 43 This has led to its usefulness as a preventive treatment being questioned. Nevertheless, fibrates are the primary indication in hypertriglyceridemia (target value <150 mg/dL).

The primary indication for niacin, which is not available in Spain, is to increase HDL concentrations, usually in combination with a statin. 44 Fibrates can also be used to raise HDL cholesterol levels, 40 although extreme care should be taken if using them in combination with a statin (fenofibrate in this case) because of the possibility of myositis. This is particularly true in polymedicated patients or those who have renal dysfunction. 45

**Antiplatelet Treatment**

Diabetes, MS, and even obesity are all prothrombotic conditions, with hyperaggregability and hypofibrinolysis. Acetylsalicylic acid has been shown to prevent atherothrombotic complications. 46 The majority of clinical practice guidelines therefore recommend doses of 81-325 mg/day, unless contraindicated, in all patients with diabetes and MS. 14,26

Figure 9 summarizes overall preventive treatment for cardiovascular and metabolic risk in patients with DM2, 26 and which can be applied almost in their entirety to patients with MS. 13

**SPECIFIC TREATMENTS**

**Hyperglycemia**

The aim of glucose regulating treatment is to reduce baseline blood glucose levels, glycohemoglobin (<7%), and postprandial blood glucose levels. There is little data available on treatment strategies in patients with MS with respect to control of the carbohydrate mechanism. The most appropriate drugs in obesity and MS are acarbose, metformin, and the glitazones, administered alone or in combination depending on the circumstances. 26

As regards DM2, the ADA/EASD treatment algorithm opts for strict control of glycohemoglobin using multiple treatments (Figure 10). 47 Several studies have demonstrated that controlling glucose values can be an effective preventive measure. 48
Weight loss is associated with relevant reductions in CVD whether DM2 is present or not. The ideal objective is to reach a BMI of 25 (a more realistic value might be <27) or a waist circumference under that indicated as the threshold in the MS criteria. As well as diet and exercise, some drugs are useful for weight reduction. To date, only sibutramin and orlistat have been approved for use, though both have poor outcomes and are only moderately well-tolerated. Rimonabant is a selective CB1 receptor blocker which has proven effective in achieving substantial weight loss and in reducing triglyceride levels and glycohemoglobin. It has also been shown to increase HDL cholesterol levels in obese patients and specifically in patients with diabetes. Its possible psychological side effects require further study.

Bariatric surgery is indicated in morbidly obese patients or, in less advanced cases, when CVD or diabetes are also present.

**Figure 9.** Overview of preventive treatment in patients with diabetes, also applicable in metabolic syndrome. ASA indicates acetylsalicylic acid; CA, calcium antagonist; ARA-II, angiotensin II receptor antagonist; Bβ, beta-blockers; ED, erectile dysfunction; CVD, cardiovascular disease; IMTC, intima-media thickness of carotid artery; HDL, high density lipoproteins; ACE, angiotensin converting enzyme; BMI, body mass index; ABI, ankle/brachial index; LDL, low density lipoproteins; MAU, microalbuminuria; BP, blood pressure; MDCT, multi-detector computed tomography; TG, triglycerides.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Ideal Target</th>
<th>Priority Indication</th>
<th>Complementary</th>
</tr>
</thead>
<tbody>
<tr>
<td>BMI/Waist</td>
<td>&lt; 25/&lt; 98 or 80 cm</td>
<td></td>
<td>Orlistat, Rimonabant</td>
</tr>
<tr>
<td>LDL</td>
<td>&lt; 70 mg/dL</td>
<td>Statin</td>
<td>Ezetimibe</td>
</tr>
<tr>
<td>HDL</td>
<td>&gt; 50 mg/dL</td>
<td></td>
<td>Niacin, Fibrato</td>
</tr>
<tr>
<td>TG</td>
<td>&gt; 150 mg/dL</td>
<td>Fibrate</td>
<td></td>
</tr>
<tr>
<td>PA</td>
<td>&lt; 130/80 mmHg</td>
<td></td>
<td>IECA/ARA-II, CA, Tiazide, β-β</td>
</tr>
<tr>
<td>Aggregation</td>
<td>?</td>
<td>ASA</td>
<td></td>
</tr>
</tbody>
</table>
Vascular Complications

Peripheral vascular complications should be managed using percutaneous or surgical procedures corresponding to the condition in question.

As regards cardio-ischemic complications, treatment should be similar to that used in any other case of chronic angina. Antianginal drugs and revascularization will be used depending on level of functioning and coronary anatomy. The classic antianginal drugs (nitrates, beta blockers, and calcium antagonists) can all be used in patients with MS, obesity, or DM2.28

In acute coronary syndromes, strict glucose control using insulin perfusion is essential and has been shown to reduce the incidence of some adverse outcomes.3 In acute cases, the recommended values are as close as possible to 110 mg/dL, and always <180 mg/dL.14 In non-critical hospitalized patients, prandial insulin can be used, with control values ranging from 90 to 130 mg/dL (fasting) and <180 mg/dL after meals.14

Criteria for revascularization in patients with DM2 are influenced by the greater risk (which requires the use of more intensive treatments) and lower success rates associated with revascularization procedures in these patients.56,57

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