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bid), fenofibrate (200 mg, qd) and omega-3 fatty acids (1 g, tid). Plasma triglycerides decreased in a few days, followed by blood cholesterol (Fig. 1). After lipid pattern improvement, insulin treatment was completely discontinued, and the patient was successfully treated with only diet and metformin; also subcutaneous heparin was stopped, while fibrates and polyenoic fatty acids were maintained. His weight decreased by 3–4 kg during the hospital stay, while 6 months afterwards his BMI was 25.2 kg/m².

In short, we report a case of a patient with massive dyslipidaemia associated with glycaemic decompensation, in the framework of previously unrecognised type 2 diabetes, anamnestic hints of “slight”, not further defined, hyperglycaemia and a familial history of diabetes. Increased HbA1c plasma levels suggest that diabetic decompensation occurred during weeks or months before, and not in a few days time, while high C-peptide plasma levels in the context of overt hyperglycaemia testify to a relative insulin deficiency and marked insulin resistance. Finally, the absence of immune markers rule out the presence of latent autoimmune diabetes in the adult (LADA), a peculiar form of smoldering type 1 diabetes that appears during adulthood. Dyslipidaemia is a common finding in diabetic populations, and represents a major coronary heart disease risk factor [1]. It may be found when type 2 diabetes is diagnosed, persisting in spite of hypoglycaemic therapy and closely related to insulin-resistance parameters. It is generally characterised by moderate hypertriglyceridaemia and low HDL-C plasma levels, while total and low-density lipoprotein-cholesterol (LDL-C) concentrations are similar to non-diabetic subjects; however LDL particles are smaller and denser than usual, and possibly more atherogenic. Moreover, the LDLs and HDLs are triglyceride-enriched [2, 3]. Less frequently, diabetes begins with a chylomicronemic syndrome. In our patient, hyperlipidaemia was at first characterised by very high triglycerides and cholesterol plasma levels. Blood triglycerides exceeding 2000 mg/dl are a rare clinical finding; when such values are present, several haematochemical parameters may look abnormal as a result of analytical interference or uneven analyte distribution (the “false” hyponatraemia we found is ascribable to the presence of sodium only in the watery plasma phase, while the lipid phase, about 10% of total plasma volume, is not permeable to sodium).

The most common primary causes of severe hypertriglyceridaemia are familial LPL deficiency and familial apo C-II deficiency (the physiological LPL activator), which may account for up to 10% of all individuals referred to a lipid clinic with the chylomicronaemia syndrome [4]. The transient hypertriglyceridaemia of our patient was not in agreement with familial conditions, characterised by rather steadily increased lipid values (also above 5–10 000 mg/dl), and not responsive to drug treatment. The coexistence of high plasma cholesterol and triglycerides with fasting chylomicrons may suggest type III hyperlipidaemia. This lipid disorder (also known as ‘dysbetalipoproteinaemia’ or ‘broad-beta disease’) is a rare condition (1:10 000 subjects) and much more frequent in subjects with an uncommon apolipoprotein E genotype (ε2/ε2). It is characterised by the development of premature atherosclerosis and plasma accumulation of partially catabolised chylomicrons and VLDLs [5]. Dyslipidaemic patients show increased cholesterol and triglyceride levels (>300 mg/dl) and a peculiar “broad beta” band between β (LDL) and pre-β (VLDL) bands at lipoprotein electrophoresis; chylomicrons may be present in variable amounts. In this case,
however, a very distinct electrophoretic β-band and a common apolipoprotein ε3ε3 genotype rule out the hypothesis of type III hyperlipoproteinaemia.

Another rare hyperchylomicronaemic syndrome (with a prevalence of about 1:10 000 in the general population) is Fredrickson type V dyslipidaemia. In this condition both VLDL and chylomicron levels are increased, while plasma triglycerides are very high. It may display either a primary or a secondary origin: in the latter case obesity, pregnancy, ethanol intake, estro-progestins, β-blockers or anion exchange resins, hypothyroidism, kidney disease and diabetes may favour its appearance in predisposed individuals. Our patient seems classifiable in this type of dyslipidaemia. Likely some concurrent lipid metabolism abnormality (e.g., familial combined hyperlipidaemia or heterozygous LPL deficiency) may represent the fuel promoting hyperchylomicronaemia (+VLDLs) in response to hyperglycaemic decompensation, most probably feed-back by incoming hypertriglyceridaemia as part of a well known vicious cycle [6, 7]. This fact might explain both transient type IIb hyperlipidaemia during clinical type V remission, and the familial history of coronary heart disease. His lipid profile 6 months after hospital discharge was alternatively normal, or suggestive of type IV hyperlipaemia. Our diagnosis is probable but the pathophysiology is presumptive, as important data are lacking (LPL activity or gene sequencing); however therapy response and patient follow-up add reliability to this interpretation.

Interestingly our patient was substantially asymptomatic: clinical complications of hyperchylomicronaemia (acute pancreatitis, eruptive xanthomatosi) are particularly common with plasma triglyceride levels above 2000 mg/dl [4], but sometimes, unpredictably, they may be absent. The relationship between hyperchylomicronaemia and premature atherosclerosis is still controversial, denied by most authors, but has received some valuable confirmation [8].

Treatment improved the lipid pattern after only a few days. A lipid-poor diet, in particular, represents the cornerstone of early treatment. Fibrates are then particularly indicated: after interacting with a specific nuclear receptor (peroxisome proliferator activated receptor-α, PPARα), they fasten LPL-induced lipolysis (especially by blocking apo C-III synthesis, a main lipoprotein lipase-inhibitor), and decrease liver triglycerides and apolipoprotein B secretion [9]. Heparin was effective in the short term, but its role in severe hypertriglyceridaemia is still under discussion [10]. It has been defined as a “plasma clearing factor” showing hypotriglyceridaemic effect in an acute setting, and may be useful when pancreatitis supervenes, especially in conjunction with insulin treatment [10, 11]. Heparin may release LPL from the endothelial surface of extrahepatic tissues, promoting its access to blood where the enzyme hydrolyses VLDL and chylomicron triglycerides. However, after a quick peak effect, chronic heparin infusion causes lipase activity to reach a plateau that is substantially lower than the former level (about 15%), decreasing correspondingly the tissue stores (likely by hepatic catabolism of circulating LPL molecules) [12]. This makes its long-term utilisation particularly questionable. Omega-3 fatty acids are another debated drug: they display hypotriglyceridaemic effect when administered to subjects with high VLDL plasma levels, while in type V patients there is no consensus on their efficacy (they could even be detrimental, favouring chylomicron synthesis in enterocytes). Also plasmapheresis may represent, in selected contexts, a useful but extreme therapeutic tool [13].

Finally, insulin effect must not be underestimated: it is always advisable in diabetic patients when marked hypertriglyceridaemia is recognised. In fact insulin can stimulate LPL production, inhibit endogenous triglyceride synthesis and promote their catabolism [14].

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D-dimer testing: advantages and limitations in emergency medicine for managing acute venous thromboembolism

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Patients with suspected deep vein thrombosis (DVT) or pulmonary embolism (PE) are frequently admitted to an Emergency Department (ED) for initial evaluation. However, management of patients with suspicion of acute venous thromboembolism (VTE) in this clinical setting can be difficult; in fact symptoms and signs of DVT are non-specific and can be found in a broad spectrum of non-thrombotic disorders. An accurate and timely objective diagnosis is necessary for immediate and correct identification of patients with acute VTE, while avoiding the bleeding risk associated with unnecessary anticoagulant therapy in patients where DVT or PE have been ruled out. The diagnostic approach to patients with suspected VTE includes clinical evaluation, diagnostic imaging and D-dimer testing [1, 2].

In a recent issue of Internal and Emergency Medicine, Siragusa [3] exhaustively discussed the currently used assays, clinical indications and limitations of D-dimer testing for managing acute VTE in emergency medicine. Measurement of D-dimer values, a degradation product of cross-linked fibrin, is a simple laboratory test, and has been extensively studied in numerous prospective cohort studies in cases of suspected DVT or PE, showing a high negative predictive value. The author [3] shows that D-dimer testing has sufficient diagnostic accuracy for ruling out acute VTE if used in combination with standardised clinical judgement. The review is important for at least three reasons.

First, many physiological and pathological conditions can increase plasma D-dimer levels (pregnancy, age, trauma, cancer, inflammation and several other clinical conditions); on the other hand D-dimer levels may fail to increase in patients with acute VTE for multiple reasons (impaired fibrinolytic activity, use of heparin or oral anticoagulants, onset of symptoms more than two weeks before blood sampling). For these reasons, D-dimer testing has a high sensitivity but a low specificity in the diagnosis of acute VTE; in clinical practice it may be associated with frequent “false positive” results, but also more rarely with “false negative” findings. However, its use should always be associated with a careful clinical assessment to rule out the presence of acute VTE only in symptomatic patients. Overuse or misuse of the D-dimer screen for VTE may have negative consequences, in terms of a burden both for patients and for healthcare costs; in fact, despite clinical guidelines, inappropriate and unnecessary measurement of D-dimer is a significant clinical problem [4].

Second, commercially available assays are very different for many reasons. Assays differ in the type of antibody used as well as in the type of data provided, as some are qualitative whereas others are quantitative. Moreover, the value of various D-dimer assays is based upon several factors including sensitivity, specificity, cost, time required, reproducibility and complexity. Finally, the diagnostic accuracy of D-dimer relies on the cut-off values, that is the range within which the test has the best sensitivity and specificity, and may be very different comparing one assay to another. Cut-off points are established by the manufacturers, according to the characteristics of D-dimer testing; a wide range of sensitivities and specificities have been reported for different D-dimer assays. Currently, no D-dimer reference standard exists, making interassay correlation poor; moreover, because of variations and differences among different assays, results from one of them cannot be extrapolated to another.

Third, another interesting issue concerning the use of the D-dimer test in emergency medicine is its use when urgent diagnostic imaging for DVT or PE is not immediately available, as for example during a night-time or weekend evaluation. Siragusa and coworkers [5] studied whether a combined approach consisting of pre-test clinical probability, D-dimer testing and low-molecular-weight heparin (LMWH) allowed deferral of objective assessment of DVT or PE for up to 72 h in 409 patients admitted to ED for suspected VTE. Pending diagnostic confirmation, patients identified with a high probability or with a moderate probability and positive D-dimer test received a protective full dose of LMWH, while the remaining patients were discharged without anticoagulant treatment. The study demonstrates the safety of this approach.

Despite the promise of these findings, there are several points of caution worth noting with respect to the use...