The Calvary After The Abort Due To Varied Attempts To Treat Rheumatoid Arthritis By Tnfa Inhibitors And The Final Resolution, Owing To Classic And Orthodox Remedies In An Old Lady Suffering From A Complex Syndrome: A Special Case Report.

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Abstract: Scope of this modest case report is to demonstrate that it is not impossible to conciliate the usage of common and orthodox medicaments to combat difficult outcomes deriving from an inconceivable series of syndroms in a single individual. The very importance I want this paper has to herald is that TNFα inhibitors may result perilous and sometimes lethal (in peculiar cases) for elder, besides to propose a treatment of that complicated syndrome by the aids of common and orthodox remedies. 

Keywords: TNFα inhibitors, INR, diabetes mellitus type 1, psoriatic rheumatoid arthritis.

I. Background

The Case Report Deals with an Ancient Lady suffering from the Following Syndromes:
- psoriatic rheumatoid arthritis
- diabetes mellitus type 1
- diverticulitis
and is 84 y. old, female and underwent 10 years ago to double knee replacements.

The volunteer (suffering from the syndrome) was not but my mother and for, I have had not to satisfy all the diktats of the Camorra imposed by almost all the Italian Local Ethical Committees, and since I am a doctor (subjected to the Galen’s oath), I avail myself of Sir Thomas Percival’s code on medical ethics(1847). Her psoriatic rheumatoid arthritis was diagnosed the very first time when the volunteer was 69 y. old. She has been proving 4 kinds of TNFα inhibitors for 12 years, (each of every drug showed a three-yearly efficacy) and this sequence of drugs had destroyed at all her immunological system (she presented always 100000 C. FUs of E. coli in urines; she had loss completely appetite; she grew progressively completely sideropenic and developed a grade 4 cardiac murmur and moreover it must be kept on account that she had never suffered from many cardiac failure throughout all her life) and when she used to take painkillers (in case of acute attacks of disabling arthritis) the assumption of these drugs forced her to stay at home because of severe risk of diarrhoeic episodes.

After this pharmacological Armageddon (euphemistically speaking) she has necessitated one entire year to:
- reset her compromised immunological system and stop her severe and recurrent urinary tract infections regain appetite fot all types of aliment present a perfect electrocardiogram with no cardiac murmur of any grade do not suffer from recurrent diarrhoeic episodes, if sometimes she assumed Nsaids. Nevertheless joint pains, due to the rheumatic arthritis, were devastating and the only remedy she could tolerate resulted prednisone (10 mg/pro day), that is known is contraindicated in case of diabetes type 1, and thus doses of insulins had to be adjusted when required. All this has implied a complex therapeutic carreau, that could never be the same or almost predictable, depending on the quality and quantity of carbohydrates she had ingested during the main meals during the day.

For one entire year this was the therapeutic programme:
- prednisone 5mg at morning and 5-7 IU of fast-acting insulin
- prednisone 5mg at 4 p.m. and 6-8 IU of fast-acting insulin
after supper 18 IU of long-acting insulin.

After one year she manifested an important DVT (deep vein thrombosis) at her right leg, with drastic difficulty of deambulation and severe thromboembolic risk at brain and lungs and her INR was 0.91. Some colleagues of mine (an haematologist and a diabetologist) have supposed that chronic usage of prednisone involves the formation of clots that drives to deep vein thrombosis,DVT (1,2) and even knee replacement can constitute a serious hazard for thromboembolic episodes in lower imbs.(3) and moreover peaks of hyperglycaemia (even if sporadic) when exceed 350 mg/dL may drive the same to thromboembolic episodes at lower extremities(4,5)

The great concern began so to come up with the possibility of coordinating the administration of Painkillers (avoiding absolutely corticosteroids)
- The insulines
- The anticoagulants.

It must be stressed that the patient, when she was under TNFα inhibitors could not take Nsaids, because of the severe diarrhoeic flows, but now, after one year she has stopped with the aforementioned drugs, she does not present the same problem and so the choice of antiinflammatory medicaments is fortunately broad:
- ketoprofene
- diclofenec
- sulindac

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- sodium naproxene
- indomethacin

at moderate or high dosages to alternate to acetaminophen.

Paying attention that some of these NSAids can alterate the glycemic value, one or two IU of fast-acting insulin had to be added (in the morning, at lunch time and at 4 o’clock in the afternoon) to the regular dosage.

II. Materials and methods

And so the schedule of painkillers and insulines became the following:

at morning: acetaminophen (1 g) and 4 IU of fast acting insulin (depending on the carbohydrates assumed at morning breakfast or no IU of insulin)

at lunch time: diclofenac (100 mg) or indomethacin (50 mg) and 8 IU of fast-acting insulin

at supper time: acetaminophen (1 g) and 16 or 18 IU of long-acting insulin, depending on carbohydrates assumed by diet.

As far as anticoagulants, in order to reach a INR comprised between 2 and 3, the schedule has been the following:

first day: coumadin (5 mg) at 4 p.m. and mesoglycane (50 mg) at supper time

from the second to the fourth day: coumadin (5 mg) at 4 p.m. and coumadin 2.5 mg at supper time.

After the fourth day the INR reached the value 1.72, I repute this value quite satisfying.

according to the protocol of the initiation of warfarin, one may behold in Table I:

<table>
<thead>
<tr>
<th>Day therapy</th>
<th>INR value</th>
<th>Total day dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>Day 1</td>
<td></td>
<td></td>
</tr>
<tr>
<td>In 2-3 days after initiation</td>
<td>&lt; 1.5</td>
<td>5-7.5 mg/die</td>
</tr>
<tr>
<td></td>
<td>1.5-1.9</td>
<td>2.5-5 mg/die</td>
</tr>
<tr>
<td></td>
<td>2.0-2.5</td>
<td>2.5 mg/die</td>
</tr>
<tr>
<td></td>
<td>&gt;2.5</td>
<td>Need for a re-check</td>
</tr>
<tr>
<td>After 4th day</td>
<td>&lt; 1.5</td>
<td>7.5-10 mg/die</td>
</tr>
<tr>
<td></td>
<td>1.5-1.9</td>
<td>5-10 mg/die</td>
</tr>
<tr>
<td></td>
<td>2.0-3.0</td>
<td>5-10 mg/die</td>
</tr>
<tr>
<td></td>
<td>&gt;3</td>
<td>Need for a re-check</td>
</tr>
</tbody>
</table>

in order to achieve the INR I had foretold.

III. Results

The craved result is achieved, so I have fain prosecuted with the suggested doses by the Protocol (6), administering coumadin 2.5 mg/die until the 8th day when I have the test of INR effectuated. Finally, the INR value was 2.2 (for this complex clinical picture, the range of INR 2-3 is more than favorable). Concomitantly, some A. A. (6), assert that corticosteroids may increase the INR value after some days in patients taking coumadin and the augmentation of the value is observable after 6.7 +/- 3.3 days from the first assumption of corticosteroids. So at 9th day the volunteer began to take again prednisone at the same prior dosage and after other 10 days the INR value was 2.8. The therapy could remain the same for long time and aught of irreparable occurred.

IV. Discussions

Inr 2.8 avoids risks of pulmonary embolism and cerebral venous thrombosis as well, as I feared at the first occurrence of symptoms of DVT in the volunteer. This case is very particular, but I deem the clinical trial and the results can be useful for everybody (especially women over 60 suffering from DVT and rheumatic arthritis evoked by diverse causes, instead of diabetes mellitus type 1).

V. Conclusions

The remission after 12 years of assumption of TNFα inhibitors has been dramatic and those drugs should be baished, especially in Italy where Camorra and Mafia of the Hospitals reigns as Sovereign.

References


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