Letters to the Editor

Body Mass Index, Cholesterol Level and Poststroke Mortality

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Olsen et al. [1] found that the body mass index (BMI) in patients with stroke was inversely related to total poststroke mortality. The mechanism underlying this association is however not clearly understood. We suggest that one possibility might be related to the higher cholesterol levels associated with high BMI.

It is well known that cholesterol levels are inversely related to all-cause mortality in middle-aged and older men and women, and a reverse causation is unlikely to account for this finding [2]. It may result from an increase in both cancer and noncardiovascular disease mortality with lower cholesterol levels.

In particular, higher serum cholesterol concentrations have been shown to be associated with lower short-term mortality after stroke [3–5]. Furthermore, Olsen et al. [6] have recently reported an inverse relation between cholesterol and stroke severity, and consequently with poststroke mortality, thus suggesting that higher cholesterol favors the development of small-vessel disease and thereby less severe strokes associated with lower mortality.

Therefore, higher levels of cholesterol might be a plausible mechanism for the inverse relationship between BMI and poststroke mortality.

References


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Reply to the Letter by Mascitelli et al.: Body Mass Index, Cholesterol Level and Poststroke Mortality

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We appreciate the thoughtful comments of Mascitelli et al. In our paper we did not adjust for serum cholesterol as this variable was not available in our database. However, for the purpose of our study adjustment for serum cholesterol is not necessary. Along with hypertension and diabetes, cholesterol is a mediator and not a confounder. Hence, these variables should not be adjusted for in the survival model, as they are part of the causal chain between BMI and death/survival. For this reason we presented two models, one including all the variables and one excluding mediators. The latter, we believe, is the most correct one. Both models, however, gave the same result: overweight persons with stroke had a better outcome than those who were underweight and normal weight. Moreover, compared to normal-weight individuals, survival was better in the obese and very obese individuals with stroke.

Whether cholesterol, as suggested by Mascitelli et al., is responsible for the inverse relationship between BMI and survival is still not clarified, we believe. The possibility exists but other unknown elements in the causal chain are possibilities as well. Our study showed a clear association between BMI, diabetes and hypertension, both risk factors associated with an increased mortality rate. So, there is no doubt that the mortality rate is increased in many overweight and obese persons. However, our study shows that this is most likely due to hypertension and diabetes and not overweight and obesity per se. It can be hypothesized that overweight and obesity signal a lifestyle, which is leading to hypertension and diabetes and thereby decreased survival. If, however, an overweight or obese person has a ‘healthy’ lifestyle, survival is not endangered.

We agree with Mascitelli et al. in their interpretation of our data. In patients with stroke the majority of poststroke deaths are caused by another stroke [1]. The risk of stroke is increased in overweight and obese persons but these strokes may be caused by small-vessel disease rather than large-vessel disease [1]. This might in part explain the lower mortality associated with overweight and obesity.

References


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