Evidence Table 98. KQ 3—Continuous Blood urea nitrogen (BUN)

| **Author Year****Study Design** | **CHD Risk Category** | **CVD drug** | **CVD****Dose (mg/d)** | **Supplement/ Control** | **N** | **Definition of outcome / Unit of outcome** | **Post treatment Mean/****(Median)****SD/SE****Lower limit (IQR, 95% CI)****Upper Limit (IQR, 95% CI)** | **Mean change/% change from baseline/****SD/****P Value** | **Between group differences in Means/ medians****SD/****P Value** | **Additional Comments** | **Overall Risk of Bias (ROB) Assessment** |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| Lee200836Parallel | At high risk for CHD | Warfarin | 3.5 | Ginseng | 12 | NRUnit: mg/dl | Mean:9.45SD: 5.16 | Mean:-2.28 | NR | There were no statistically significant differences between the two groups. | N/A |
| No Treatment | 13 | Mean:13.46SD: 4.44 | Mean:0.46 |
| Napoli199852Parallel | Unclear | Pravastatin | 20-40 | Vitamin E |  | NRUnit: NR | NR | NR | NR | No differences in routine laboratory tests or adverse events were registered during the study. In particular there were no differences between the values recorded in each individual after treatment compared to baseline values, and no differences between the treatment groups. | Medium |
| No Treatment |  | NR | NR |