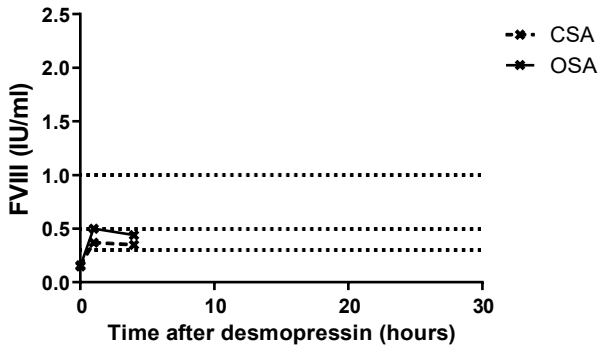
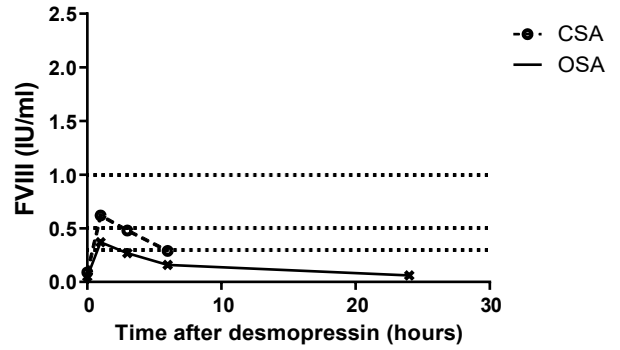


Supplementary Data 4 – Individual desmopressin administrations

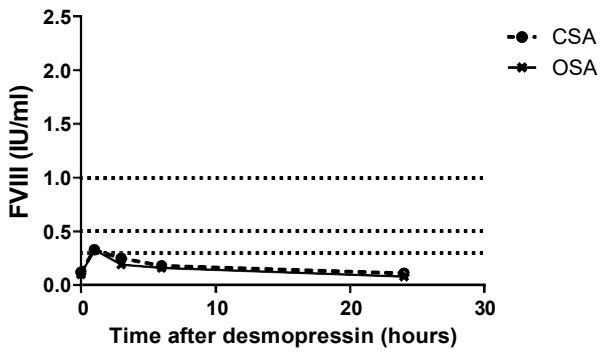
9



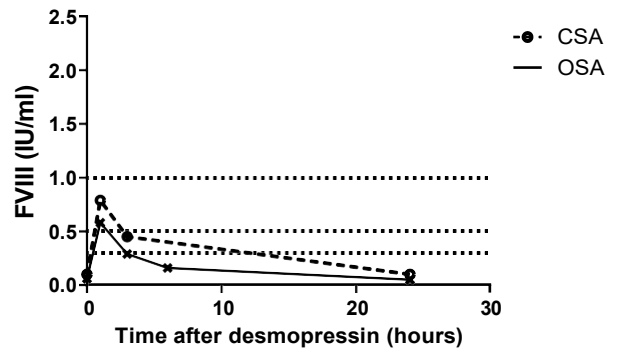
108



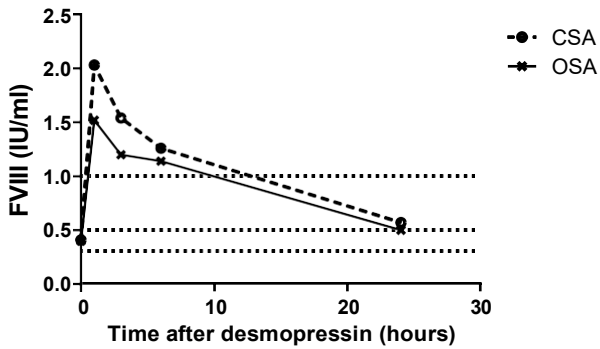
109



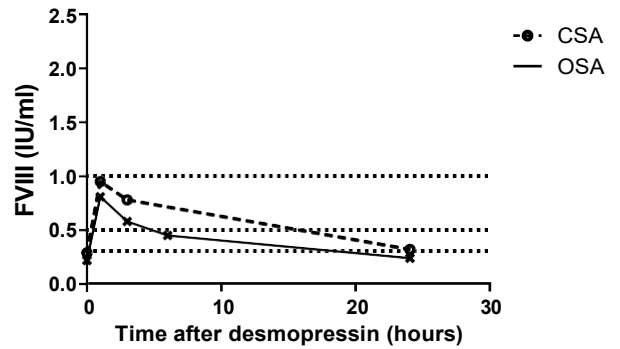
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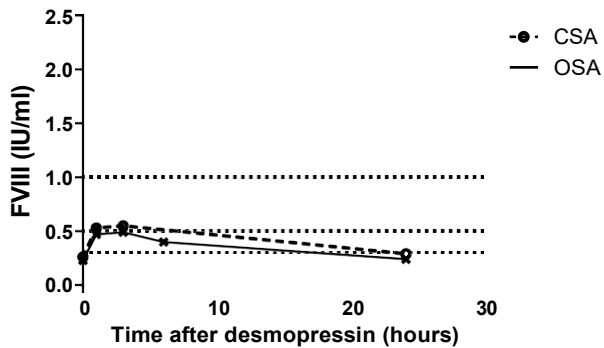
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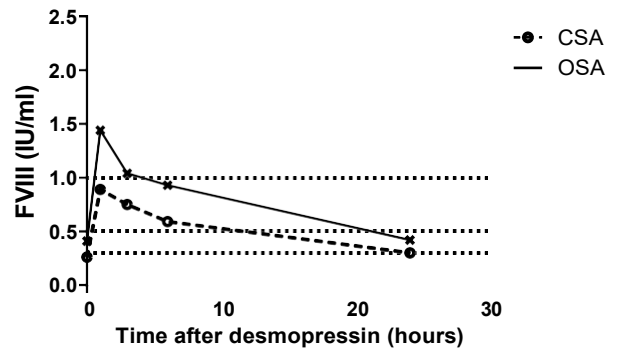
112



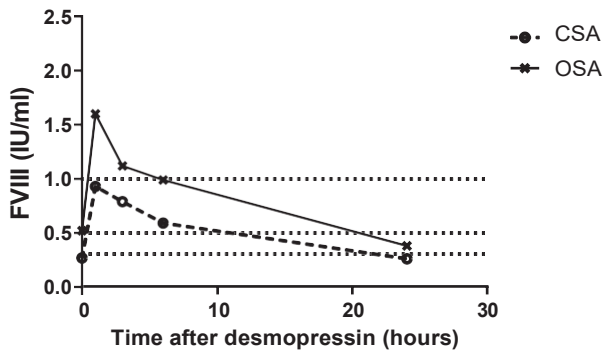
114



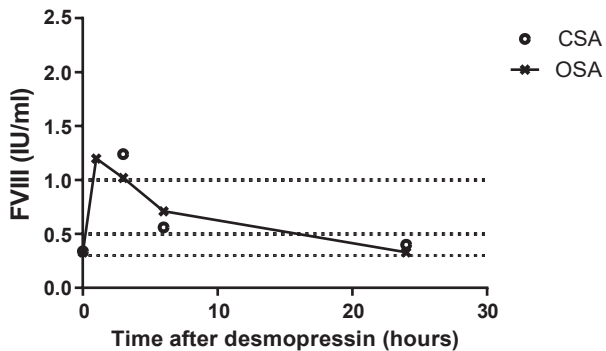
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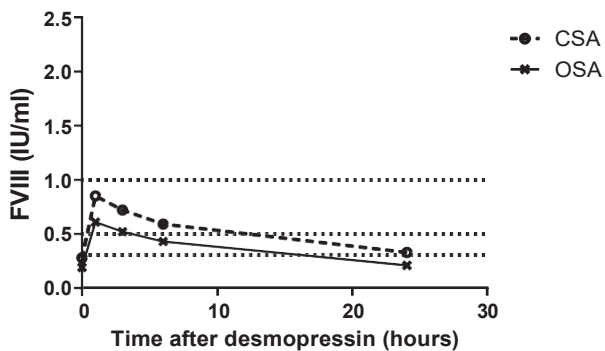
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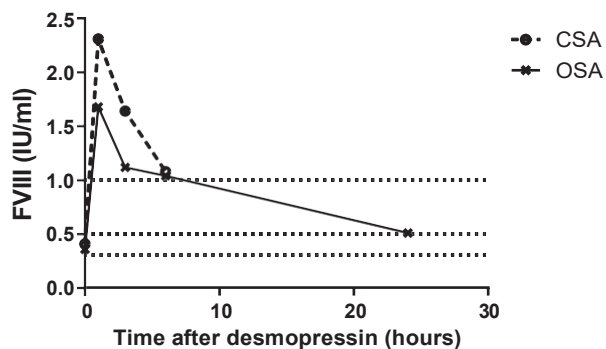
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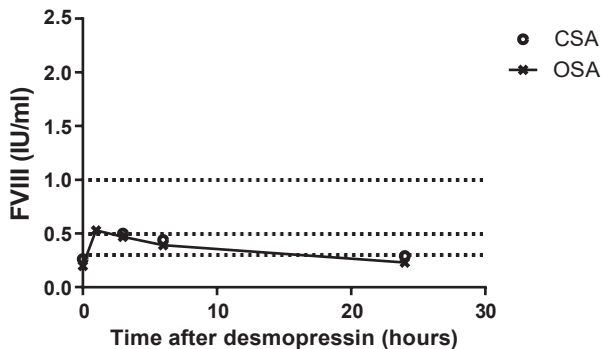
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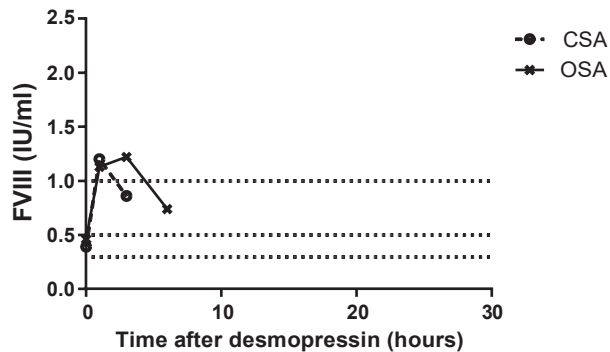
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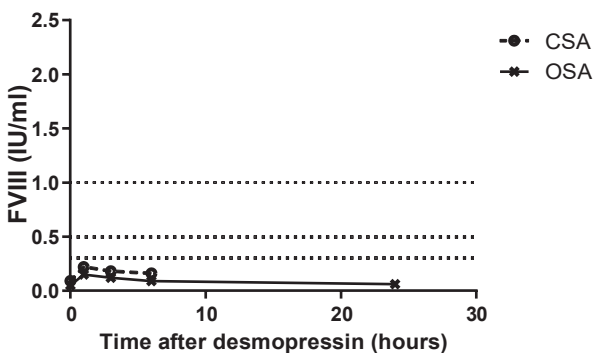
122



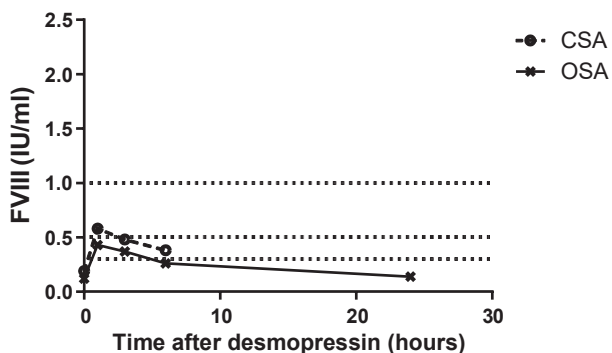
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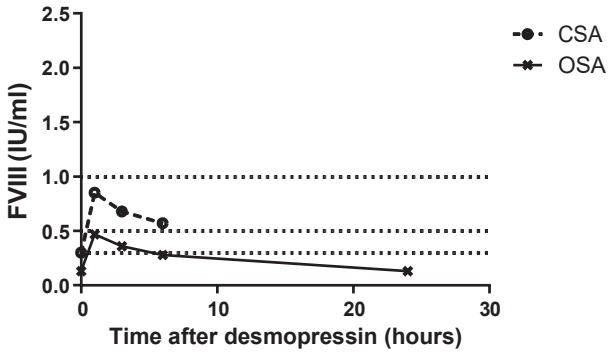
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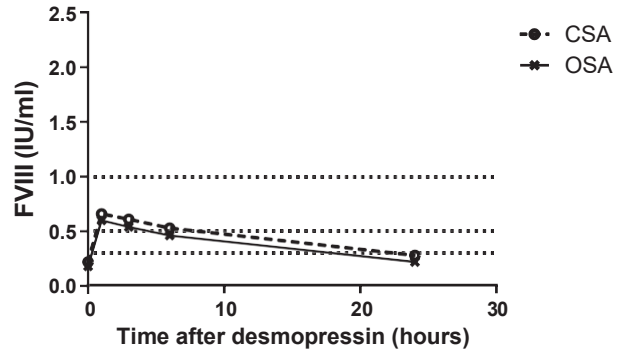
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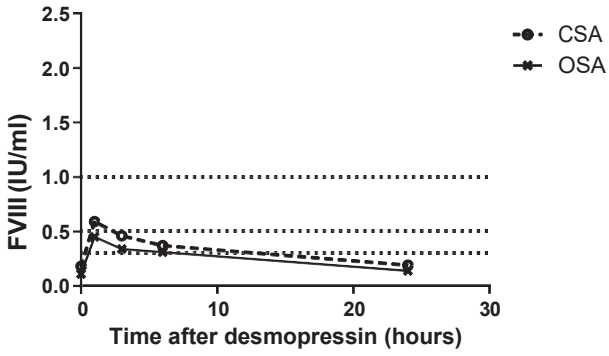
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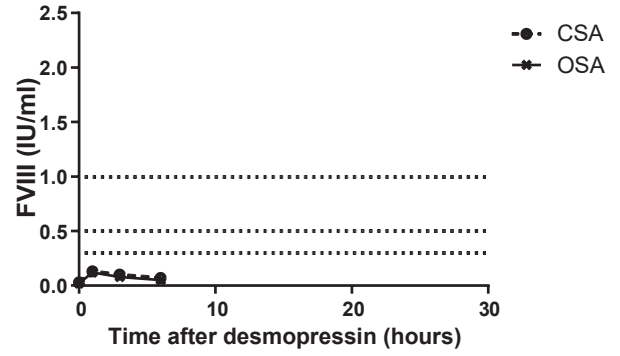
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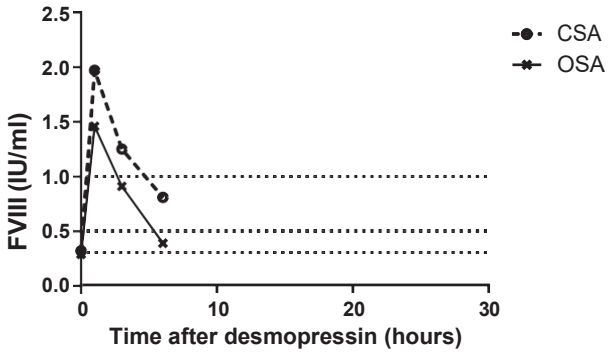
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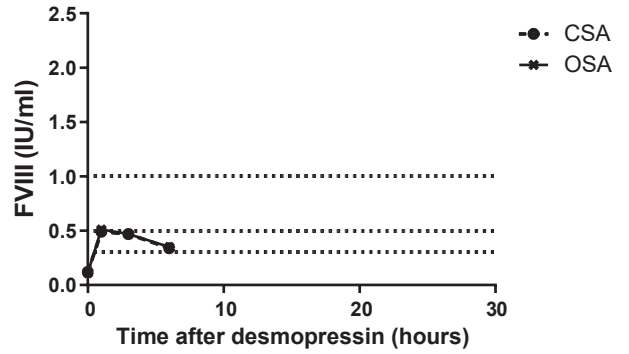
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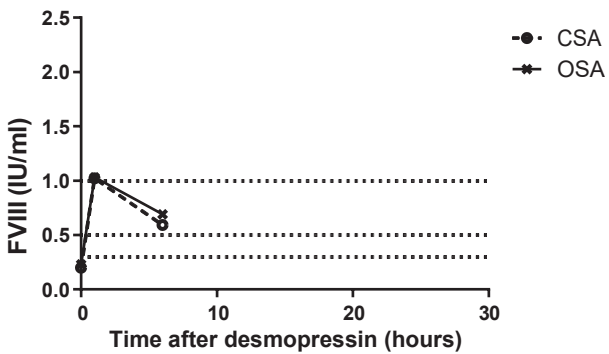
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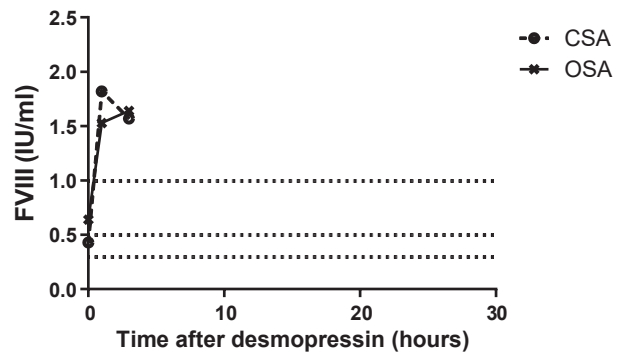
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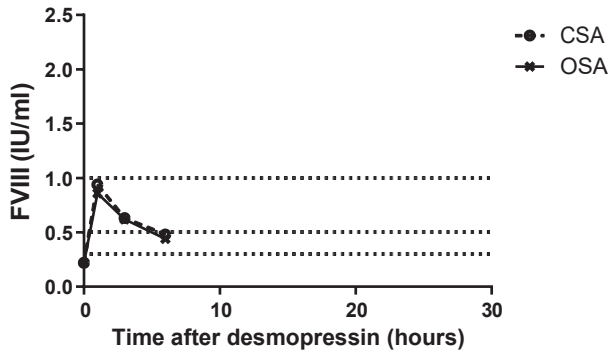
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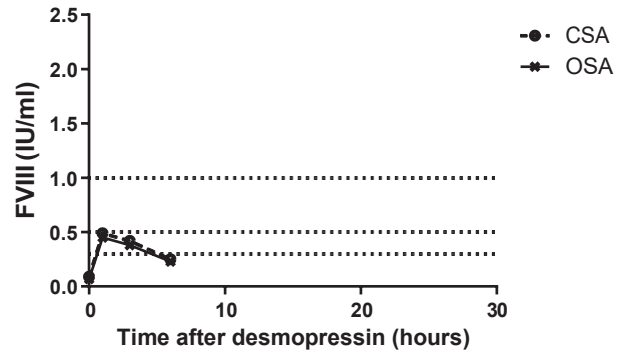
143



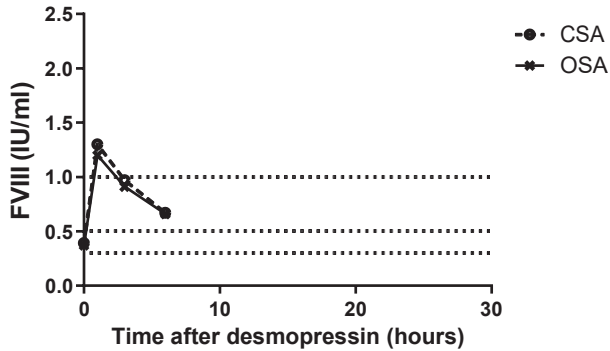
144



145



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Individual FVIII measurements of all 27 patients receiving desmopressin measured by the one-stage assay (OSA; solid lines) and by the chromogenic assay (CSA; dashed lines). Lines were not depicted if time points were not consecutive. T0 is before desmopressin, all other time points are after the end of desmopressin administration. Dotted horizontal lines represent partial response (PR, 0.30 IU/ml), complete response (CR, 0.50 IU/ml) and 1.0 IU/ml.