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## Sialic acid O-acetylation as characteristic marker for recombinant erythropoietins

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### Abstract

The presence of O-acetylations on sialic acids of recombinant erythropoietins (EPO) has been described in previous literature [1,2]. Shahrokh *et al.* observed that Dynepo (epoetin delta), an epoetin produced in a human cell line but no longer marketed, contained no O-acetylations in contrast to three other originator erythropoiesis stimulating agents (Eprex, NeoRecormon, Aranesp)[3]. In 2011, Reichel presented first high accuracy mass spectrometric data of human urinary erythropoietin at the MDI workshop. Human urinary EPO did also not show O-acetylations (for sensitivity reasons only O-glycans were investigated). In order to clarify whether O-acetylations of sialic acids may pose a specific marker for recombinant erythropoietins, a comprehensive mass spectrometric study was performed specifically on the O-glycans of 40 EPO pharmaceuticals including many biosimilar epoetins.

For more details refer to the full article published in Drug Testing and Analysis [4].

### References:

- [1] Rush RS, Derby PL, Smith DM, Merry C, Rogers G, Rohde MF, Katta V (1995) Microheterogeneity of erythropoietin carbohydrate structure. *Anal Chem.* **67**:1442-52.
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- [3] Shahrokh Z, Royle L, Saldova R, Bones J, Abrahams JL, Artemenko NV, Flatman S, Davies M, Baycroft A, Sehgal S, Heartlein MW, Harvey DJ, Rudd PM. (2011) Erythropoietin produced in a human cell line (Dynepo) has significant differences in glycosylation compared with erythropoietins produced in CHO cell lines. *Mol Pharm.* **8**:286-96.
- [4] Reichel, C. (2013), Differences in sialic acid O-acetylation between human urinary and recombinant erythropoietins: a possible mass spectrometric marker for doping control. *Drug Test Analysis*, **5**: 877-889.