

WADA Technical Letter - TL14

Document Number:	TL14	Version Number:	3.0
Written by:	WADA Science		
		Approved by:	WADA Executive Committee
Reviewed by:	WADA Laboratory Expert Group		
Date:	21 December 2020	Effective Date:	1 January 2021

DIFFERENCE IN "A" AND "B" SAMPLE URINE CHARACTERISTICS

WADA wishes to draw the attention of the <u>Laboratories</u> to the following issue that may affect <u>Laboratory</u> operations. This pertains, in particular, to significant differences in urine characteristics, related to color and/or turbidity, between urine "A" and "B" *Samples*, which may be observed upon inspection at *Sample* receipt.

Upon reception, *Samples* shall not be rejected for analysis based on noted color/turbidity differences between the "A" and "B" *Samples*. Should such differences between the "A" and "B" *Samples* be observed, the following steps shall be conducted by the <u>Laboratory</u> in collaboration with the associated <u>Testing</u> Authority:

• Before opening the "A" bottle, the <u>Laboratory</u> shall separately document the observed difference(s) and inform the *Testing* Authority in writing as soon as possible:

[Comment: It is recommended that the correspondence includes a picture of the "A" and "B" Samples (as provided in the appendix below).]

[Comment: A single freeze/thaw cycle of both "A" and "B" Samples may address the difference in turbidity; therefore, storing the "A" and "B" Samples frozen overnight and then thawing them may be an option before reporting the differences to the <u>Testing Authority</u>. It seems that this procedure, in some cases, may result in urines with identical turbidity].

• The <u>Laboratory</u> shall proceed to analyze the "A" *Sample*, for the requested test menu, bearing in mind to preserve as much "A" *Sample* volume as possible for additional investigation, if necessary:

[Comment: The <u>Testing Authority</u> and the <u>Laboratory</u> shall bear in mind the availability of the "B" Sample, which may be split for further investigations, if necessary.]

- The <u>Testing Authority</u> (or <u>Results Management Authority</u>, if different) should investigate with the <u>Doping Control</u> Officers (DCOs) whether any observations or unusual circumstances occurred during the <u>Sample</u> collection procedure. In case of abnormality in the collection procedure, the <u>Testing Authority</u> should evaluate other <u>Samples</u> collected from the <u>Athlete</u> and, in addition, the <u>Athlete</u> should be targeted for further <u>Sample</u> collections;
- In the case of a <u>Presumptive Adverse Analytical Finding (PAAF)</u>, the finding shall be confirmed and the result reported in *ADAMS* accordingly. The difference between the "A" and "B" <u>Samples</u> and any other relevant observation shall be recorded in the comments section of the Test Report. The <u>Testing Authority</u> (or <u>Results Management Authority</u>, if different) shall conduct <u>Results Management</u> of the case taking into account the <u>Laboratory</u> analytical results and observations, and determine the next steps including the "B" <u>Sample</u> confirmation;
- The Testing Authority (or Results Management Authority, if different) shall alert the associated

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<u>Athlete Passport Management Unit</u> (<u>APMU</u>) to investigate the *Athlete*'s steroid profile with particular attention to this specific *Sample*.

Examples of color/turbidity differences between "A" and "B" Samples





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