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W. Schänzer, H.
Geyer, A. Gotzmann,
U. Mareck-Engelke
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und Buch Strauß,
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321–326. Google
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To avoid such tedious
and lengthy
procedures, vacuum
matrix-assisted laser
desorption ionization

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(vMALDI) coupled
with the linear ion
trap mass
spectrometry
(LIT/MS) technique is
tested for its
applicability as a
rapid screening
technique.

Commonly used
doping agents like
nandrolone,
boldenone,
trenbolone,

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testosterone, and betamethasone were chosen as study compounds.

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steps. Initially,
screening of samples
is performed, in the
case of a suspicious
result; an additional
selective
confirmation is
carried out.[16]

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Gas chromatography

mass spectrometry

(GC MS) has been

the traditional

technique for doping

control analysis of

anabolic steroids in

biological samples.

Although liquid chro

matography mass

spectrometry (LC/MS)

has become an

important technique

in doping control, the

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detection of
In
saturated
hydroxysteroids by
LC-MS remains a
problem due to their
low ionization
efficiency under
electrospray.

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data to be used to
detect designer
steroids.

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Preventive doping
control analysis:
liquid and gas ...

Nowadays sensitive
LC-MS applications
are available for
almost all classes of
doping agents but
more research is

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needed for the detection of peptide hormones. New developments in column particle size ($3 \mu\text{m}$) allow separations on shorter columns, and 50 mm columns are becoming the standard in doping analysis.

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respect to scan
speed, sensitivity as
well as polarity
switching allow an
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Erythropoietins
(EPOs) are substances
listed in S2 of the

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Prohibited List and
are used commonly
by athletes to
increase endurance
performance.
According to the
current WADA
Technical
Documents, sarcosyl-
polyacrylamide gel
electrophoresis (SAR-
PAGE) followed by

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western blotting to
differentiate
erythropoietins
based on their
molecular weights is
the only method that
can be used for both
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types of
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This study presents a fast multi-analyte screening method specifically developed for the detection of xenobiotics in urine. The proposed method allows the screening of several classes of substance in a single

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method with a run-
time of 11 min,
inclusive of post-run
and reconditioning
times.

Chromatographic
separation is
achieved in 7.2 min
using a reversed-
phase 2.7 μm fused-
core particle column,
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exceeding 400 bar
and therefore
enabling the use ...

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“ Blood Doping
Screening through
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Electrophoresis ” C.
Harrison (San Diego
State University,
California, USA) To
facilitate the
continued anti-
doping efforts it is
becoming important
to make the best use
of the limited analysis

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The growth-based
rapid microbiology
testing segment is
expected to account
for the largest share
of the market in

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2018. The large share of this segment can be attributed to factors such as the ease of processing (this method uses conventional liquid or agar media), the limited requirement of skilled professionals, and supportive government regulations.

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