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procedures such as
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separation,

pediatric heart and
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pancreas and small
intestine

transplantation,

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procedures, peroral

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myotomy, jugular

foramen tumors,

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surgeries, proton

beam therapy and

many other

complex surgeries

and procedures.

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Uncommon and

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by anesthesia

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and APRN/PA
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middle- and low-
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policymakers on
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investment policies
for new industrial
development
strategies.

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printed edition of
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Vitamin K-
Dependent Proteins
in Relation to
Human Health" that
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is intended to
support the design,
implementation,
analysis,
interpretation, and
quality evaluation of
registries created
to increase
understanding of
patient outcomes.

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condition, or Egd

exposure, and that
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serves one or more
predetermined

scientific, clinical,
or policy purposes.

A registry database
is a file (or files)
derived from the
registry. Although
registries can serve
many purposes, this
guide focuses on
registries created

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for one or more of the following purposes: to describe the natural history of disease, to determine clinical effectiveness or cost-effectiveness of health care products and services, to measure or monitor safety and harm, and/or to measure

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quality of care. Egd

Registries are classified according to how their populations are defined. For example, product registries include patients who have been exposed to biopharmaceutical products or medical devices. Health services registries

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consist of patients
who have had a
common procedure,
clinical encounter,
or hospitalization.

Disease or
condition registries
are defined by
patients having the
same diagnosis,
such as cystic
fibrosis or heart
failure. The User ' s
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by researchers
affiliated with
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(Developing
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Effectiveness)
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were subject to

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multiple internal
and external
independent
reviews.

This volume
provides a
comprehensive and
world-class review
of the field of
histiocytic
neoplasms and
hemophagocytic
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(HLH). It reviews

all the advances in
the field of

histiocytoses during
the last ten years,

particularly with
regards to the

genomic findings in
LCH and other

histiocytic

neoplasms and the
new suggested

classification of the
histiocytic

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disorders. 2014 Egd

Additionally, it features a state-of-the-art update on the most recent treatment strategies for LCH, including the results of the last LCH-III international trial, salvage therapies such as reduced-intensity conditioning (RIC)

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stem cell transplant (SCT), and targeted therapies with BRAF and MEK inhibitors, as well as the challenging cases of CNS-neurodegenerative LCH and its therapeutic perspectives. For primary and secondary HLH the book updates the

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pathophysiological findings, including macrophage-activation syndrome (MAS), and

includes a special chapter on HLH in adults. Treatment chapters encompass therapy for newly diagnosed HLH and refractory disease

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as well as stem-cell transplantation and novel therapies.

The text also highlights the most recent advances in the treatment of the uncommon histiocytic disorders, such as Erdheim-Chester disease (ECD), Juvenile xanthogranuloma

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(JXG) and JXG-like conditions, Rosai-Dorfman disease (RDD), and the very rare malignant histiocytoses.

Written by international experts in the field, Histiocytic Disorders is a valuable resource for clinicians, researchers,

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fellows and residents who are interested in or manage histiocytic disorders in children and adults.

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represent an

opportunity to treat

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the whole field of
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imaging, diagnostic investigation of cyst fluid, and those using pancreatic juice; and also examines aspects ranging from the development of malignancy to the timing and method of resection, focusing on both

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