Wilms Tumor Educational & Global Resource Library

Wilms Cancer Foundation Launches Global Multilingual Educational Library for Wilms Tumor and Childhood Kidney Cancer

New international educational platform provides multilingual Wilms tumor information for children, parents, caregivers, survivors, healthcare professionals and researchers worldwide

FOR IMMEDIATE RELEASE

WASHINGTON, DC, USA - August 18th, 2026 - The **Wilms Cancer Foundation (WCF)** has announced the launch of its **Global Educational Library**, a growing international collection of multilingual educational resources about **Wilms tumor, pediatric renal cancer and childhood kidney cancer**.

The Global Educational Library has been created to improve worldwide access to clear, reliable and practical information covering the complete Wilms tumor journey—from symptoms and diagnosis through treatment, relapse, survivorship and long-term follow-up care.

Resources are being developed for children, parents, caregivers, survivors, healthcare professionals, researchers, students, advocates and organizations working to improve childhood cancer outcomes internationally.

A central focus of the initiative is the production and translation of Wilms tumor educational materials into multiple languages. By making specialist childhood kidney cancer information available in the languages used by different communities, WCF aims to reduce language barriers, strengthen health literacy and improve access to knowledge in both well-resourced and underserved healthcare settings.

A Global Educational Library Dedicated to Wilms Tumor

Wilms tumor, also known as nephroblastoma, is the most common kidney cancer diagnosed in children. Although it is highly treatable in many countries, survival outcomes continue to vary considerably according to geography, early diagnosis, specialist expertise, access to treatment, healthcare infrastructure and the availability of reliable information.

Families searching for information about childhood kidney cancer may encounter medical terminology, fragmented resources or materials that are unavailable in their primary language. Healthcare professionals in countries with limited pediatric oncology resources may also have restricted access to specialist educational materials.

The WCF Global Educational Library has been developed to help address these challenges by bringing Wilms tumor information together within one accessible international educational platform.

The library will include:

- Plain-language Wilms tumor guides for parents and caregivers
- Childhood kidney cancer fact sheets and downloadable resources
- Wilms tumor briefing documents and professional educational materials
- Information about symptoms and the early signs of Wilms tumor
- Guidance explaining Wilms tumor diagnosis, staging and risk classification
- Educational resources covering surgery, chemotherapy and radiation therapy

- Information about relapsed and refractory Wilms tumor
- Nutrition guidance for children undergoing cancer treatment
- Wilms tumor survivorship and long-term follow-up information
- Resources addressing the late effects of childhood cancer treatment
- Global reports examining treatment access and survival inequalities
- Clinical and educational toolkits for healthcare professionals
- Visual resources, maps, infographics and caregiver quick guides
- Materials supporting Wilms tumor research and international collaboration

New resources, translations and language editions will be added progressively as the Global Educational Library continues to expand.

Multilingual Wilms Tumor Resources for Families Worldwide

The production of multilingual childhood cancer information is an important part of WCF's international education strategy.

Language can become a significant barrier when a child is diagnosed with cancer. Parents and caregivers may be required to understand unfamiliar medical terminology, treatment plans, potential side effects and follow-up requirements while experiencing considerable emotional distress.

Providing educational information in a family's preferred language can help them:

- Understand what Wilms tumor is
- Recognize the possible symptoms of childhood kidney cancer
- Prepare for diagnostic tests and staging
- Understand surgery, chemotherapy and radiation treatment
- Identify questions to discuss with their child's medical team
- Learn about nutrition and supportive care during treatment
- Understand relapse, surveillance and follow-up care
- Access information about survivorship and long-term effects

WCF's multilingual educational work is intended to complement—not replace—the guidance provided by qualified healthcare professionals and each child's treating medical team.

Supporting Healthcare Professionals and Childhood Cancer Organizations

The Global Educational Library is also designed to support doctors, nurses, surgeons, pediatric oncologists, allied health professionals, researchers, medical students and childhood cancer organizations.

Its resources can help healthcare and nonprofit communities strengthen:

- Wilms tumor awareness and early recognition
- Parent and caregiver education
- Communication between families and treatment teams
- Healthcare professional education and knowledge sharing
- Nutrition and supportive-care guidance
- Survivorship education and long-term follow-up
- International research collaboration
- Childhood cancer advocacy and health policy
- Adaptation of educational materials for different countries and communities

By making reusable educational resources available internationally, WCF aims to help organizations provide locally relevant information without requiring every healthcare system or childhood cancer group to create materials independently.

Addressing Global Inequalities in Childhood Kidney Cancer Care

Children diagnosed with Wilms tumor do not experience equal outcomes worldwide.

In many lower-resource settings, delayed diagnosis, limited pathology and imaging services, shortages of essential medicines, restricted access to pediatric cancer specialists, treatment abandonment and the absence of family education can reduce the likelihood of successful treatment.

The Global Educational Library will support wider efforts to improve:

- Awareness of Wilms tumor symptoms
- Earlier childhood kidney cancer diagnosis
- Access to understandable treatment information
- Education for parents and caregivers
- Knowledge sharing between healthcare professionals
- Survivorship and long-term follow-up care
- International collaboration and advocacy
- Access to resources in low- and middle-income countries

The initiative aligns with the wider international objective of reducing inequalities in childhood cancer survival and supports the principles of the **World Health Organization Global Initiative for Childhood Cancer**.

Building Wilms Tumor Topical Knowledge Across the Complete Care Pathway

The Global Educational Library forms part of WCF's broader educational ecosystem and its commitment to developing comprehensive knowledge around every major area of Wilms tumor and childhood kidney cancer.

Content will connect users with information covering:

- What Wilms tumor is and how it develops
- Wilms tumor causes, genetics and risk factors
- Symptoms and early warning signs
- Diagnostic imaging, biopsy and pathology
- Wilms tumor stages and risk groups
- Surgery and nephrectomy
- Chemotherapy drugs and side effects
- Radiation therapy
- Stage 4 and metastatic Wilms tumor
- Bilateral Wilms tumor
- Relapsed Wilms tumor
- Nutrition during childhood cancer treatment
- Emotional and practical family support
- Treatment completion and surveillance
- Long-term effects and survivorship
- Childhood kidney cancer research and clinical trials
- Global treatment access and survival outcomes

Connecting these subjects within a structured international library will help parents, professionals and researchers move more easily between related information and develop a more complete understanding of Wilms tumor.

Part of the Global Wilms Tumor Initiative™

The Global Educational Library supports the objectives of the **Global Wilms Tumor Initiative™**, WCF's international framework for improving awareness, education, advocacy, earlier diagnosis, survivorship and global collaboration.

Through this initiative, WCF is working to:

- Increase international awareness of Wilms tumor
- Support earlier recognition and diagnosis
- Expand access to reliable educational information
- Strengthen global advocacy for childhood kidney cancer
- Improve survivorship knowledge and long-term outcomes
- Encourage international research and healthcare collaboration

The library also complements the Foundation's wider digital and educational resources, including the **Complete Guide to Wilms Tumor**, **Wilms Tumor Knowledge Index™**, **Wilms Tumor WebApp™**, **Wilms Tumor AI Support Service™** and its international Educational Literature Programme.

Statement from the Wilms Cancer Foundation

"Every child and family affected by Wilms tumor should be able to access clear and understandable information, regardless of where they live or which language they speak," said **TJ Hodgkinson, Founder and Chief Executive Officer of the Wilms Cancer Foundation**.

"The launch of the WCF Global Educational Library represents an important step toward creating a more equitable international knowledge environment for Wilms tumor and childhood kidney cancer.

"Parents should not be prevented from understanding their child's diagnosis because suitable information is unavailable in their language. Healthcare professionals should also be able to access practical educational resources that can support communication, knowledge sharing and family care.

"By producing materials in multiple languages and making them available internationally, we are helping to connect families, healthcare professionals, researchers and childhood cancer organizations with the knowledge they need. The library will continue to grow as we develop new resources, expand into additional languages and work with partners and communities around the world."

International Collaboration and Knowledge Sharing

WCF will continue working with healthcare professionals, translators, researchers, childhood cancer organizations, patient advocates and international partners to develop, review and distribute future educational materials.

The Foundation's approach combines specialist Wilms tumor knowledge with plain-language communication, multilingual production and international collaboration.

Where appropriate, materials will be adapted to reflect regional circumstances, healthcare environments and the educational needs of different communities. This will help ensure that translated resources are understandable, culturally relevant and useful within the settings for which they are intended.

Benefits for Parents, Professionals, Researchers and Policymakers

For Parents and Caregivers

The library provides understandable information that can help families learn about Wilms tumor symptoms, diagnosis, treatment, side effects, nutrition, relapse, survivorship and follow-up care.

For Healthcare Professionals

Resources can support patient education, professional learning, multidisciplinary communication and the sharing of Wilms tumor knowledge across countries and healthcare systems.

For Researchers and Students

The library will provide structured access to educational materials, global reports, research-related resources and information about childhood kidney cancer priorities.

For Childhood Cancer Organizations

Multilingual and adaptable resources can support local awareness, family education, advocacy and community engagement.

For Governments and Policymakers

The library will help demonstrate the importance of early diagnosis, accessible information, pediatric oncology capacity, equitable treatment and long-term childhood cancer care.

Looking Ahead

The Wilms Cancer Foundation will continue to expand the Global Educational Library through:

- The publication of new Wilms tumor educational resources
- Translation into additional languages
- Collaboration with international healthcare professionals and organizations
- Development of country- and region-specific resources
- New materials for parents, caregivers and survivors
- Additional clinical and professional education
- Digital, visual and mobile-accessible learning materials
- Greater coverage of global childhood cancer inequalities
- Ongoing development of the Wilms Tumor Knowledge Index™

The long-term objective is to establish a comprehensive international knowledge resource capable of supporting everyone affected by Wilms tumor—from the moment symptoms are first identified through treatment, recovery, survivorship and long-term care.

About the Wilms Cancer Foundation

The Wilms Cancer Foundation (WCF) is an international charitable organization dedicated to improving outcomes for children affected by Wilms tumor through education, awareness, advocacy, support services, research collaboration, and global partnerships.

As an official partner of the World Health Organization (WHO), the Foundation supports the objectives of the WHO Global Initiative for Childhood Cancer (GICC) through ongoing educational, advocacy, and support initiatives designed to improve childhood cancer outcomes worldwide.

The Foundation's flagship **Global Wilms Tumor Initiative™ (GWTI)** seeks to strengthen awareness, education, support services, healthcare engagement, and international collaboration to improve the lives of children and families affected by Wilms tumor around the world.

Access the WCF Global Educational Library

Explore multilingual Wilms tumor information, childhood kidney cancer resources, caregiver guides and healthcare professional materials at:

https://www.wilmsfoundation.org/educational-literature-on-wilms-tumor

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