

Journée de formation « médecin » et « infirmier »
Journée de Recherche du GFRUP

Usage(s) du Point of Care - Viraux et infectieux

POC ? POCT ? TDR ? TROD ?

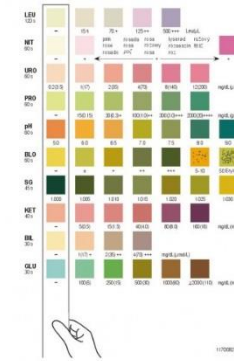
- Point-Of-Care (Test) & Test de Diagnostic rapide
 - Tests de laboratoire
 - Fabriqués sous forme permettant sa réalisation au lit du malade
 - Avec résultats immédiats (<30 mn/1h)
 - Par des personnels non biologistes ou technicien de biologie

=> Capacité à modifier la prise en charge

POC non infectieux

Multiples, en médecine générale et/ou urgence polyvalentes

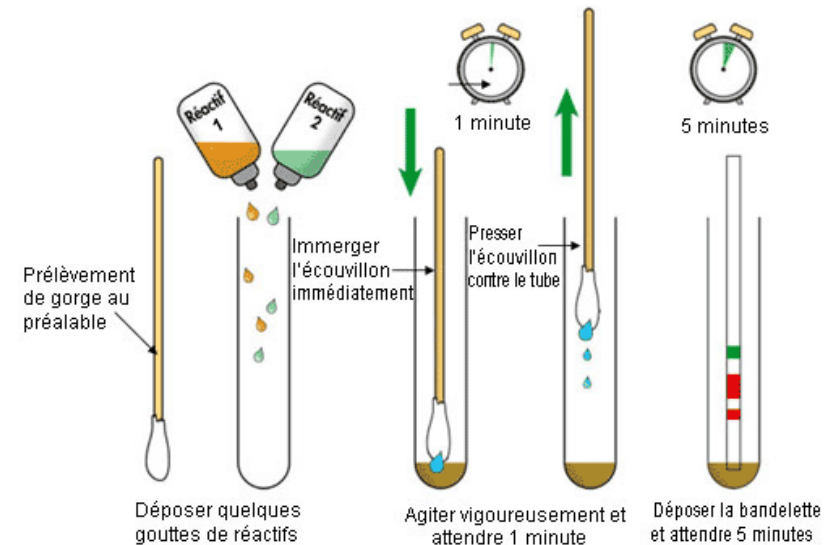
- Dextro
- HbA1C
- HCG
- INR/TP
- Ethanol
- Hemoccue
- GdS
- BU ...



Strepto-Test



- « StreptoTest »
- Recherche d'antigènes du streptocoque bêta hémolytique du groupe A
- Dans les angines :
 - orientation étiologique difficile
 - Sanction thérapeutique
 - Epargne antibiotique



CRP Capillaire



TDR Grippe

TDR VRS



Intérêt : Epargne antibiotique

Point-of-care C-reactive protein testing to reduce inappropriate use of antibiotics for non-severe acute respiratory infections in Vietnamese primary health care: a randomised controlled trial

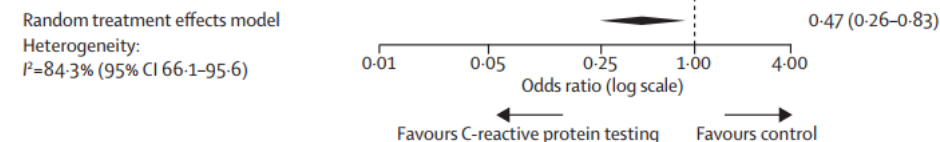


Nga T T Do, Ngan T D Ta, Ninh T H Tran, Hung M Than, Bich T N Vu, Long B Hoang, H Rogier van Doorn, Dung T V Vu, Jochen W L Cals, Arjun Chandna, Yoel Lubell, Behzad Nadjm, Guy Thwaites, Marcel Wolbers, Kinh V Nguyen, Heiman F L Wertheim



	CRP	Control	OR (95% CI)	p value
Immediate antibiotic prescription				
All patients	441/1017 (43.4%)	647/1019 (63.5%)	0.41 (0.34-0.49)	<0.0001
Children	227/510 (44.5%)	333/518 (64.3%)	0.39 (0.30-0.52)	<0.0001
Adults	214/507 (42.2%)	314/501 (62.7%)	0.40 (0.30-0.52)	<0.0001

Site	Control	CRP	OR (95% CI)
Ba Trieu	49/107 (46%)	60/110 (55%)	1.42 (0.83-2.42)
Ba Vi	82/97 (85%)	40/86 (47%)	0.16 (0.08-0.32)
Dong Da	77/83 (93%)	71/80 (89%)	0.61 (0.21-1.81)
Ha Dong	108/135 (80%)	94/129 (73%)	0.67 (0.38-1.19)
Hoan Kiem	65/81 (80%)	62/81 (77%)	0.80 (0.38-1.70)
Linh Nam	62/75 (83%)	51/72 (71%)	0.51 (0.23-1.12)
Long Bien	71/91 (78%)	49/83 (59%)	0.41 (0.21-0.79)
Mai Huong	62/92 (67%)	29/91 (32%)	0.23 (0.12-0.42)
Sai Dong	93/97 (96%)	50/80 (62%)	0.07 (0.02-0.22)
Thanh Xuan	69/89 (78%)	75/90 (83%)	1.45 (0.69-3.05)





Intérêt : Optimisation de la prise en charge

Ann Biol Clin (Paris), 2018 Oct 1;76(5):545-552. doi: 10.1684/abc.2018.1378.

Evaluation of the capillary assay of C-reactive protein (CRP) through the length of consultation in pediatric emergencies and its economic impact.

Roulliaud M¹, Pereira B², Cosme J³, Mourgues C², Sarret C¹, Sapin V³, Caron N¹, Bouvier D³.

antibiothérapie) et biologique (valeurs de CRP), l'intérêt de la CRP délocalisée a été évaluée. Dans le groupe CRP rapide, versus le groupe contrôle, une baisse significative sur les médianes du temps de consultation aux urgences (60 (IQR 33-125) versus 180 (IQR 158-208) minutes), du nombre d'actes de biologie par patient (1 (IQR 1-3) versus 7 (IQR 3-8)), du coût des examens complémentaires facturés par patient (5,4 (IQR 5,4-32,6) versus 153,8 (IQR 46,9-180,4) euros), et du coût des réactifs dépensés par le laboratoire par patient (5,2 (IQR 5,2-6,4) versus 33,2 (IQR 2,3-34,2) euros). Ainsi, dans le cadre

Roulliaud M, Pereira B, Cosme J, Mourgues C, Sarret C, Sapin V, Caron N, Bouvier D. Evaluation of the capillary assay of C-reactive protein (CRP) through the length of consultation in pediatric emergencies and its economic impact. *Ann Biol Clin (Paris)*. 2018 Oct 1;76(5):545-552

C-Reactive Protein Bedside Testing in Febrile Children Lowers Length of Stay at the Emergency Department

Ruud G. Nijman, MD,* Henriëtte A. Moll, MD, PhD,* Yvonne Vergouwe, PhD,† Yolanda B. de Rijke, PhD,‡ and Rianne Oostenbrink, MD, PhD*

Results: The preimplementation cohort included 609 children of whom 286 (47%) had traditional CRP. The postimplementation cohort included the following 1330 children: 728 (55%) children had bedside CRP and 156 (12%) children had traditional CRP. Bedside CRP significantly lowered the median LOS of children in whom an additional diagnostic CRP test was performed, from 178 minutes (interquartile range, 135–232 minutes) to 148 minutes (interquartile range, 108–200 minutes) (30 minutes, 19% of total LOS). A significant reduction of 15% of the (log)LOS remained after adjusting for other determinants of (log)LOS; propensity score analysis showed a 16% reduction.

Nijman RG, Moll HA, Vergouwe Y, de Rijke YB, Oostenbrink R. C-Reactive Protein Bedside Testing in Febrile Children Lowers Length of Stay at the Emergency Department. *Pediatr Emerg Care*. 2015 Sep;31(9):633-9



TDR Grippe

Les TDR Grippe (RIDT), un long chemin ...

Rationnel :

- Epargne examens complémentaires inutiles
- Diminue durée de séjour
- Tri des patients aux urgences
- Diminue hospitalisations inutiles
- Permet traitement précoce ...



Advantages and Disadvantages of RIDTs

Advantages

- Produce quick result in less than approximately 15 minutes, simple to perform
- Some RIDTs are cleared for office/bedside use. RIDTs that have been CLIA waived can be used in settings that include point-of-care.

Disadvantages

- Sub-optimal test sensitivity, false negative results are common, especially when influenza activity is high
- Sensitivity of RIDTs to detect influenza B viral antigens is lower than for detection of influenza A viral antigens.
- Although specificity is high, false positive results can also occur, especially during times when influenza activity is low.
- Some RIDTs distinguish between influenza A or B viruses while others do not. RIDTs that provide results on the type of influenza virus (e.g., influenza A or B virus), do not provide information on influenza A virus subtype [e.g., A(H1N1)pdm09 versus A(H3N2)] or specific virus strain information (e.g., degree of similarity to vaccine strains). RIDTs cannot distinguish between seasonal influenza A virus infection and novel influenza A virus infection (due to infection with avian or variant influenza A viruses).
- Sensitivities of RIDTs are generally approximately 50-70%, but a range of 10-80% has been reported compared to viral culture or RT-PCR. Specificities of RIDTs are approximately 90-95% (range 85-100%). Thus false negative results occur more commonly than false positive results.



Les TDR Grippe (RIDT), un long chemin ...

TDR Grippe

Impact of rapid influenza diagnostic test on physician estimation of viral infection probability in paediatric emergency department during epidemic period

Sylvie Lacroix^{a,*}, Bénédicte Vrignaud^a, Estelle Avril^a, Anne Moreau-Klein^b, Marianne Coste^b, Elise Launay^{c,d}, Christelle Gras-Le Guen^{a,c,d}

Journal of Clinical Virology 72 (2015) 141–145

The RIDT used was Quikvue influenza[®] test (Quidel Corporation, San Diego, California). The test is based on an immunochromatography technique and can be easily performed at the bedside, with results available within 10 min. All the nasopharyngeal swabs were tested with RIDT and with laboratory test: viral culture-direct fluorescent antigen and RT-PCR (gold standard). The laboratory test was considered positive if one laboratory test indicated influenza. Rodriguez et al. reported in 2002 the performance characteristics of Quikvue influenza[®] test in pediatrics. His sensitivity was 73% and his specificity was 95–99%.

Table 3
Impact of RIDT on prescriptions.

	Post clinical examination N= 170 (%)	Post RIDT N= 170 (%)	Post RIDT+ N= 80 (%)	Post RIDT- N= 90 (%)
Laboratory tests	133 (78.2)	70 (41.1) [*]	6 (7.5) ^{**}	64 (71.1)
Chest radiographs	113 (66.4)	78 (45.8) [*]	20 (25.0) ^{**}	58 (64.4)
Urine dipstick	112 (65.8)	53 (31.2) [*]	6 (7.5) ^{**}	47 (52.2)
Antibiotic		37 (21.7)	4 (5.0)	33 (36.6)
Oseltamivir	2 (1.1)	4 (2.3)	4 (5.0)	0 (0.0)
Hospitalization	39 (22.9)	29 (17.0)	11 (13.7)	18 (20.0)

^{*} p < 0.05 versus total population after the clinical examination.
^{**} p < 0.0005 versus total population after the clinical examination.



QuickVue[®]
Influenza A+B TEST

TDR Grippe

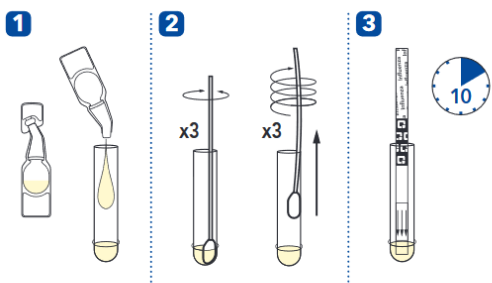


RIDT V1.0 versus RIDT V2.0 (=> DIA)

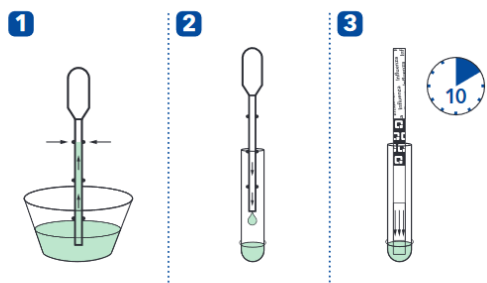


Procedure Card

Nasal Swab Procedure

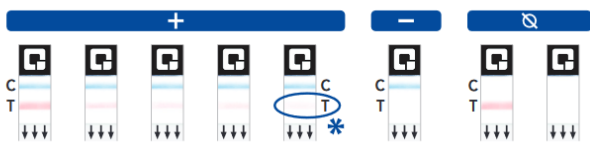


Nasal Wash / Aspirate Procedure



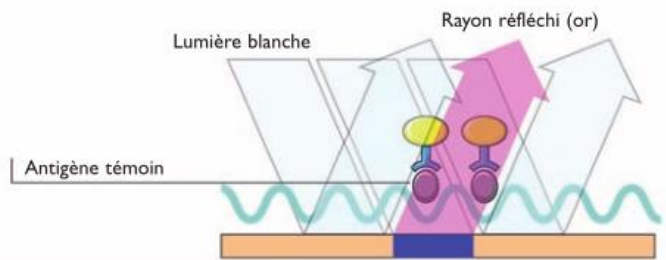
* Look closely! This is a positive result. Even if you see a very faint, pink Test Line and a blue Control Line, you must report the results as POSITIVE. The positive test line is usually very prominent, but test line intensity can vary.

C = Control Line T = Test Line



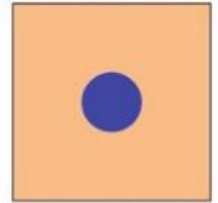
Traditional RIDT (1st generation)

NÉGATIF

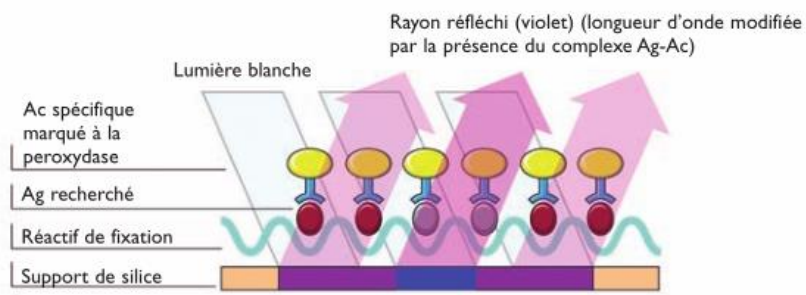


Lecture du test

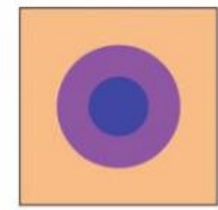
Spot central: témoin interne de la réaction



POSITIF



Spot central: témoin interne de la réaction
halo violet: réaction positive



DIA (RIDT 2nd generation)



Les TDR Grippe (RIDT), un long chemin ...

TDR Grippe

Annals of Internal Medicine

REVIEW

Diagnostic Accuracy of Novel and Traditional Rapid Tests for Influenza Infection Compared With Reverse Transcriptase Polymerase Chain Reaction

A Systematic Review and Meta-analysis

Joanna Merckx, MD, MSc; Rehab Wali, BSc, MBBS; Ian Schiller, MSc; Chelsea Caya, MScPH; Genevieve C. Gore, MLIS; Caroline Chartrand, MD, MSc; Nandini Dendukuri, PhD; and Jesse Papenburg, MD, MSc

Index Test Type

Index Test Type	Influenza A		Influenza B	
	Pooled Sensitivity (95% CrI), %	Pooled Specificity (95% CrI), %	Pooled Sensitivity (95% CrI), %	Pooled Specificity (95% CrI), %
Study population (age)‡				
Traditional RIDTs				
Children (31 influenza A studies; 9 influenza B studies)	61.2 (55.0 to 67.2)	99.2 (98.5 to 99.7)	65.7 (45.3 to 80.5)	99.6 (99.2 to 99.8)
Adults (23 influenza A studies; 5 influenza B studies)	42.6 (34.8 to 50.9)	99.5 (98.6 to 99.8)	33.2 (19.9 to 50.7)	99.9 (99.4 to 100)
Difference in RIDT sensitivity: children vs. adults	18.5 (8.4 to 28.3)	-	31.8 (6.1 to 52.6)	-
DIAs				
Children (11 influenza A studies; 11 influenza B studies)	87.6 (81.8 to 92.2)	98.1 (96.4 to 99.1)	82.5 (71.2 to 90.2)	98.8 (95.6 to 99.7)
Adults (8 influenza A studies; 7 influenza B studies)	75.4 (66.6 to 82.6)	96.7 (94.7 to 98.0)	57.0 (39.5 to 71.6)	98.8 (97.5 to 99.5)
Difference in DIA sensitivity: children vs. adults	12.1 (3.1 to 22.1)	-	25.3 (6.9 to 44.7)	-
Rapid NAATs				
Children (4 influenza A studies; 4 influenza B studies)	90.2 (79.2 to 95.8)	99.0 (96.8 to 99.8)	95.9 (82.9 to 99.2)	99.5 (98.2 to 99.9)
Adults (4 influenza A studies; 4 influenza B studies)	87.4 (71.1 to 95.6)	98.0 (93.2 to 99.5)	75.7 (51.8 to 90.7)	99.3 (97.8 to 99.8)
Difference in NAAT sensitivity: children vs. adults	2.7 (-10.7 to 19.7)	-	19.5 (1.0 to 43.7)	-



Les TDR Grippe (RIDT), un long chemin ...

TDR Grippe

Medico-economic impact of the bedside diagnosis of influenza in the paediatric emergency ward

Etude monocentrique CHU de St Etienne, **514** enfants, comparés à une cohorte rétrospective des 3 dernières années + déclaratif intention post examen clinique

11 faux positifs, 10 faux négatifs :

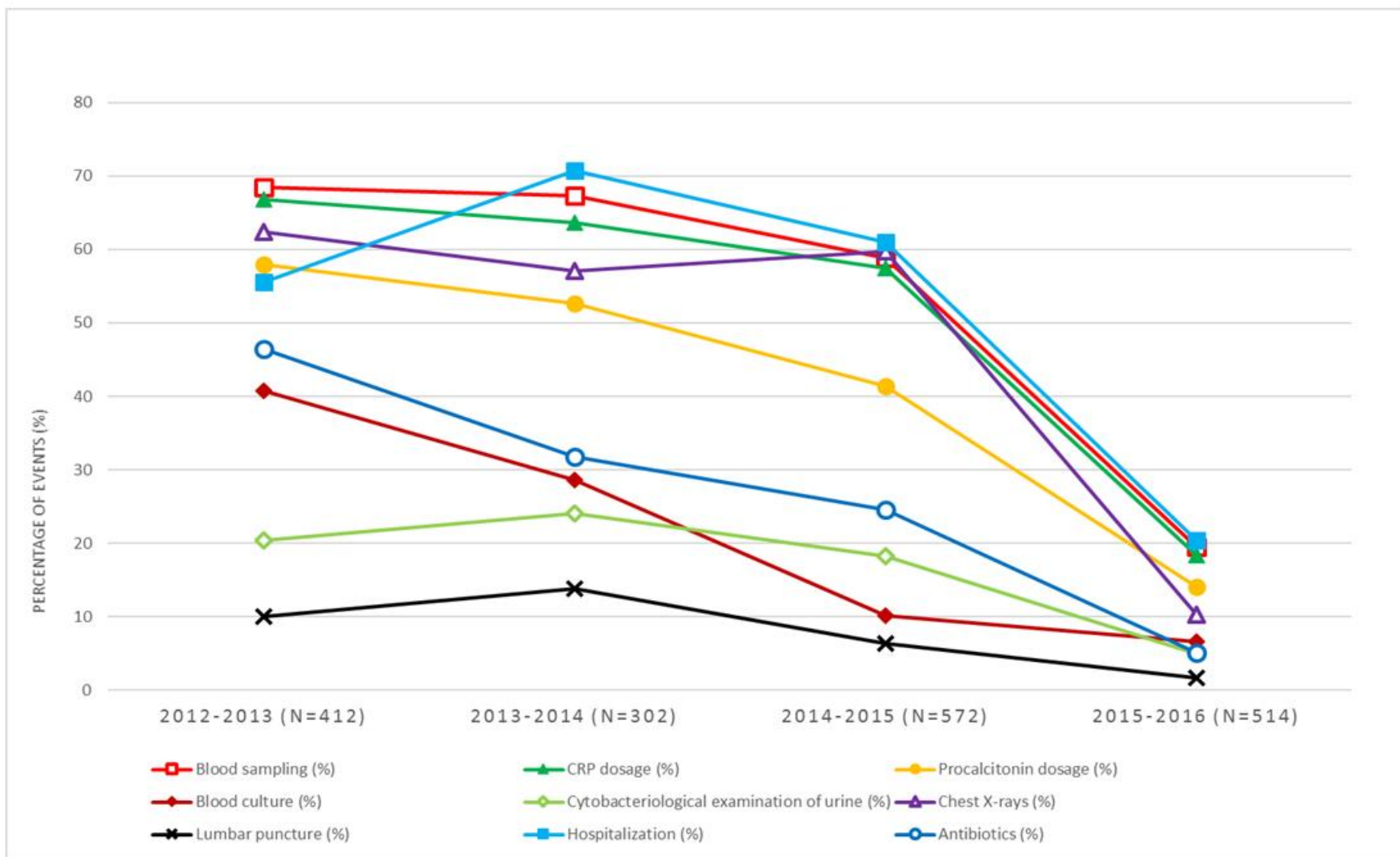
- Sensibilité 95,6 %
- Spécificité 95,2 %



Les TDR Grippe (RIDT), un long chemin ...

Medico-economic impact of the bedside diagnosis of influenza in the paediatric emergency ward

TDR Grippe





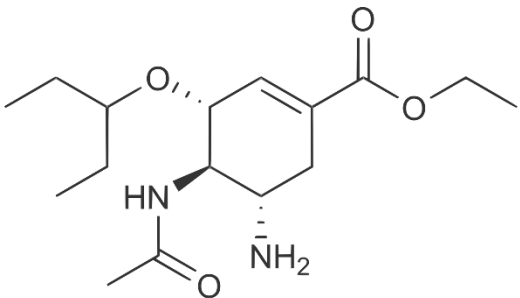
Les TDR Grippe (RIDT), un long chemin ...

Medico-economic impact of the bedside diagnosis of influenza in the paediatric emergency ward

TDR Grippe



Usage du tamiflu hyperprécoce avec TDR ?





TDR VRS

Quel intérêt ?



Bronchiolite

=

VRS

=

Pas de traitement spécifique

80 %

Bamberger E, Srugo I, Abu Raya B, Segal E, Chaim B, Kassis I, Kugelman A, Miron D. What is the clinical relevance of respiratory syncytial virus bronchiolitis?: findings from a multi-center, prospective study. Eur J Clin Microbiol Infect Dis. 2012 Dec;31(12):3323-30. doi: 10.1007/s10096-012-1699-2. Epub 2012 Jul 24.



Quel intérêt ?

Impact potentiel de l'identification du virus ?

⇒ Cohorting probabiliste chez les enfants hospitalisés

- Regrouper les bronchiolites VRS dans un secteur de soins distinct des bronchiolites « non VRS » afin d'éviter une « deuxième bronchio nosocomiale » aux enfants qui étaient VRS négatifs.
- On protège les enfants VRS neg (20%)

Problématique :

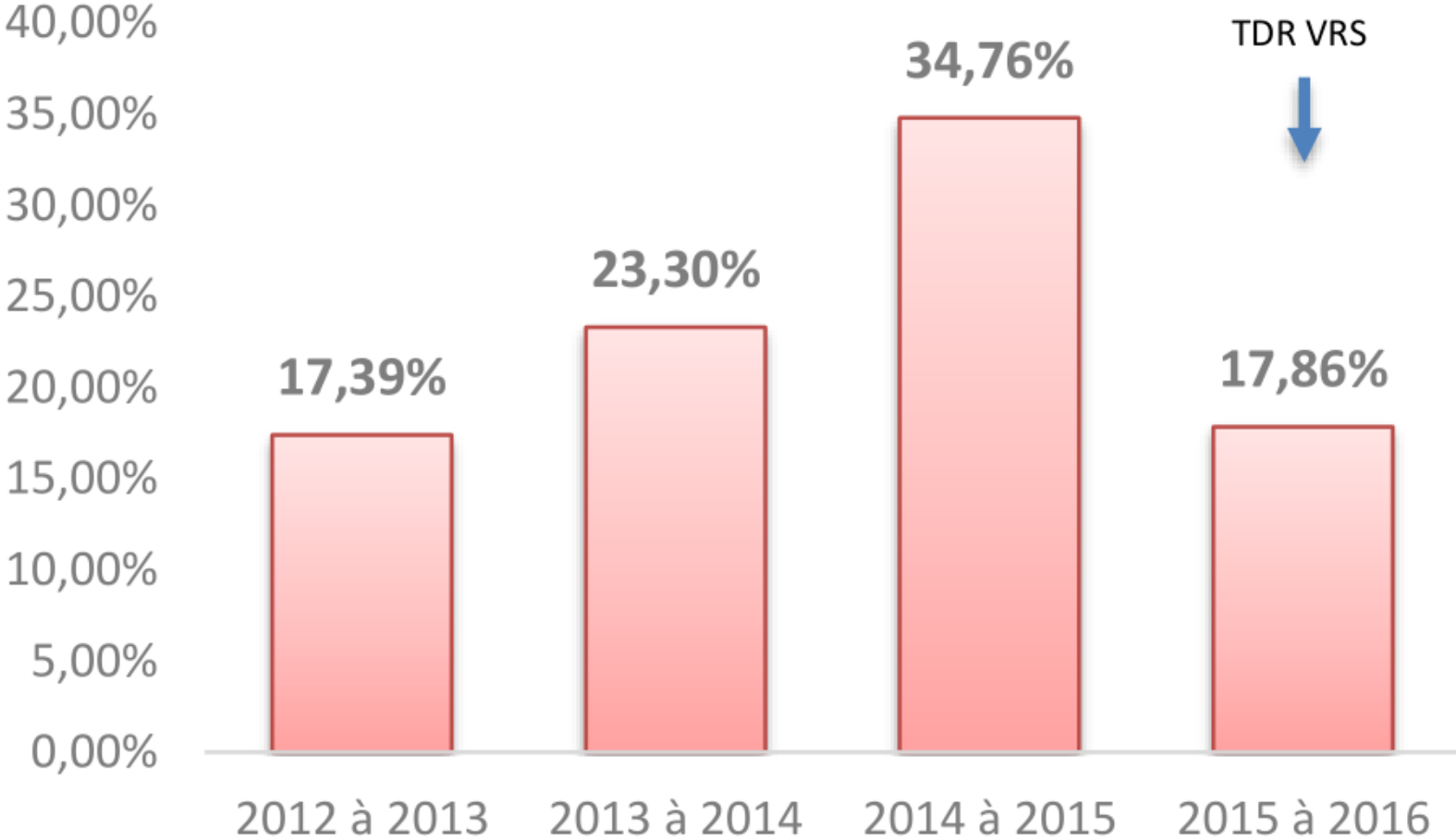
- Réponse au test viral dès les urgences avant hospitalisation en service

⇒ POC



TDR VRS

Quel intérêt ?



Les sensibilité, spécificité, valeur prédictive négative et valeur prédictive positive ont été respectivement de 97,6 ; 90,3 ; 87,5 et 98,2%.

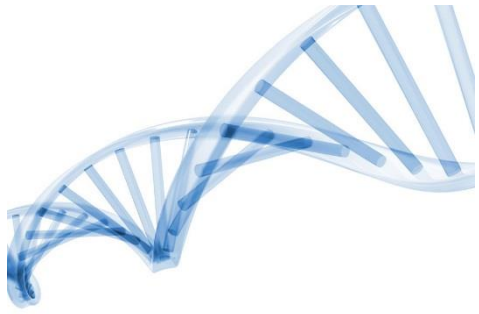
CRP Capillaire

TDR Grippe



TDR VRS

PCR POC ?



PCR POC ?

- Plus de la recherche dans certains centres
 - Se/Sp parfaite (gold standard)
 - Multiplexe
 - Systèmes POC
 - Pas d'entretien
 - Pas de personnel dédié
 - Systèmes automatisés
 - Résultats <1h
- Coûts ...

